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(73) Proprietor: **MERCK FROSST CANADA INC.**
16711 Trans-Canada Highway
Kirkland
Quebec H9H 3L1 (CA)

(72) Inventor: **Prasit, Petpiboon**
117 Argyle Drive
Kirkland,
Quebec, H9H 5A6 (CA)
Inventor: **Fortin, Rejean**

11673 Hurteau
Montreal North,
Quebec, H1J 3W8 (CA)
Inventor: **Hutchinson, John H.**
3455 Cote des Neige
No. 304 Montreal,
Quebec, H3H 1T6 (CA)
Inventor: **Belley, Michel L.**
2005 Dutrisac
No. 314 St. Laurent,
Quebec, H4L 4K8 (CA)
Inventor: **Leger, Serge**
51 Rue Lamarche
Dollard des Ormeaux,
Quebec, H9B 3E5 (CA)
Inventor: **Gillard, John**
710 Westchester Avenue
Baie D'Urfe,
Quebec (CA)
Inventor: **Frenette, Richard**
1915 De Limbource
Laval,
Quebec, H7H 4S7 (CA)

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⑦A Representative: **Cole, William Gwyn et al**
European Patent Department
Merck & Co., Inc.
Terlings Park
Eastwick Road
Harlow
Essex CM20 2QR (GB)

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Description

BACKGROUND OF THE INVENTION

5 European Patent Applications 166,591 and 275,667 disclose a series of indole-based compounds with activity as prostaglandin antagonists and inhibitors of leukotriene biosynthesis respectively. In EP 181,568 and EP 200,101 are disclosed a series of compounds, containing two aromatic nuclei, which are described as possessing activity as lipoxygenase inhibitors. In EP 279,263 is disclosed a series of indoles, benzofurans and benzothiophenes which are described as possessing activity as lipoxygenase inhibitors. U.S. 10 Patent 4,629,733 describes novel indolinones which are antithrombotic and inhibit both phosphodiesterase and tumor metastasis. The chemical preparation of quinolyndoles is referred to by Sheinkman, *et al.*, Chem. Ab., Vol. 67, 54017 (1967), without mentioning any utility for such compounds. A number of N-acyl derivatives of indole-3-acetic acid are described as potential anti-inflammatory agents by Biniecki, *et al.*, Chem. Ab., Vol. 98, 197936 (1983), by Pakula, *et al.*, Chem. Ab., Vol. 105, 190835 (1986), and in British Pat. 15 Spec. 1,228,848.

SUMMARY OF THE INVENTION

20 The present invention relates to compounds having activity as leukotriene biosynthesis inhibitors, to methods for their preparation, and to methods and pharmaceutical formulations for using these compounds in mammals (especially humans).

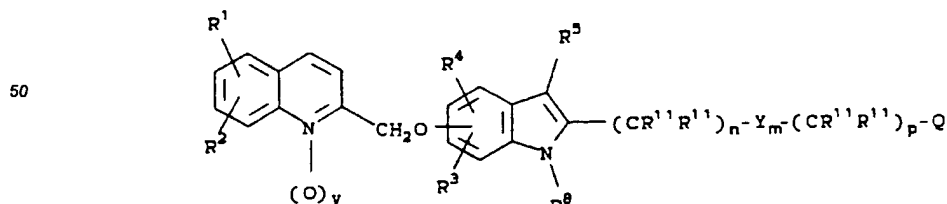
Because of their activity as leukotriene biosynthesis inhibitors, the compounds of the present invention are useful as anti-asthmatic, anti-allergic, and anti-inflammatory agents and are useful in treating allergic rhinitis and chronic bronchitis and for amelioration of skin diseases like psoriasis and atopic eczema. These 25 compounds are also useful to inhibit the pathologic actions of leukotrienes on the cardiovascular and vascular systems for example, actions such as result in angina or endotoxin shock. The compounds of the present invention are useful in the treatment of inflammatory and allergic diseases of the eye, including allergic conjunctivitis. The compounds are also useful as cytoprotective agents and for the treatment of migraine headache.

30 Thus, the compounds of the present invention may also be used to treat or prevent mammalian (especially, human) disease states such as erosive gastritis; erosive esophagitis; inflammatory bowel disease; ethanol-induced hemorrhagic erosions; hepatic ischemia; noxious agent-induced damage or necrosis of hepatic, pancreatic, renal, or myocardial tissue; liver parenchymal damage caused by hepatotoxic agents such as CCl₄ and D-galactosamine; ischemic renal failure; disease-induced hepatic damage; bile salt 35 induced pancreatic or gastric damage; trauma- or stress-induced cell damage; and glycerol-induced renal failure.

The compounds of this invention are inhibitors of the biosynthesis of 5-lipoxygenase metabolites of arachidonic acid, such as 5-HPETE, 5-HETE and the leukotrienes. Leukotrienes B₄, C₄, D₄ and E₄ are known to contribute to various disease conditions such as asthma, psoriasis, pain, ulcers and systemic 40 anaphylaxis. Thus inhibition of the synthesis of such compounds will alleviate these and other leukotriene-related disease states.

DETAILED DESCRIPTION OF THE INVENTION

45 The present invention provides novel compounds of the formula I:



I

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wherein:

- R¹, R², R³, R⁴ and R¹⁰ are independently hydrogen, halogen, lower alkyl, lower alkenyl, lower alkynyl, -CF₃, -CN, -NO₂, -N₃, -C(OH)R¹¹R¹¹, -CO₂R¹², -SR¹⁴, -S(O)R¹⁴, -S(O)₂R¹⁴, -S(O)₂NR¹⁵R¹⁵, -OR¹⁵, -NR¹⁵R¹⁵, -C(O)R¹⁶ or -(CH₂)_rR²¹;
- 5 R⁵ is hydrogen, -CH₃, CF₃, -C(O)H, X¹-R⁶ or X²-R⁷;
- R⁶ and R⁹ are independently alkyl, alkenyl, -(CH₂)_uPh(R¹⁰)₂ or -(CH₂)_uTh(R¹⁰)₂;
- R⁷ is -CF₃ or R⁶;
- R⁸ is hydrogen or X³-R⁹;
- each R¹¹ is independently hydrogen or lower alkyl, or two R¹¹'s on same carbon atom are joined to form a
- 10 cycloalkyl ring of 3 to 6 carbon atoms;
- R¹² is hydrogen, lower alkyl or -CH₂R²¹;
- R¹³ is lower alkyl or -(CH₂)_rR²¹;
- R¹⁴ is -CF₃ or R¹³;
- R¹⁵ is hydrogen, -C(O)R¹⁶, R¹³, or two R¹⁵'s on the same nitrogen may be joined to form a monocyclic
- 15 heterocyclic ring of 4 to 6 atoms containing up to 2 heteroatoms chosen from O, S or N;
- R¹⁶ is hydrogen, -CF₃, lower alkyl, lower alkenyl, lower alkynyl or -(CH₂)_rR²¹;
- R¹⁷ is -(CH₂)_s-C(R¹⁸R¹⁸)-(CH₂)_s-R¹⁹ or -CH₂C(O)NR¹⁵R¹⁵;
- R¹⁸ is hydrogen or lower alkyl;
- R¹⁹ is a) a monocyclic or bicyclic heterocyclic ring containing from 3 to 9 nuclear carbon atoms and 1 or 2
- 20 nuclear hetero-atoms selected from N, S or O and with each ring in the heterocyclic radical being formed of 5 or 6 atoms, or b) the radical W-R²⁰;
- R²⁰ is alkyl or -C(O)R²³;
- R²¹ is phenyl substituted with 1 or 2 R²² groups;
- R²² is hydrogen, halogen, lower alkyl, lower alkoxy, lower alkylthio, lower alkylsulfonyl, lower alkylcarbonyl,
- 25 -CF₃, -CN, -NO₂ or -N₃;
- R²³ is alkyl, cycloalkyl, or monocyclic monoheterocyclic ring;
- R²⁴ is the residual structure of a standard amino acid, or R¹⁸ and R²⁴ attached to the same N can cyclize to form a proline residue;
- m is 0 to 1;
- 30 n is 0 to 3;
- p is 1 to 3 when m is 1;
- p is 0 to 3 when m is 0;
- r is 0 to 2;
- s is 0 to 3;
- 35 t is 0 to 2;
- u is 0 to 3;
- v is 0 or 1;
- W is O, S or NR¹⁵;
- X¹ is O, or NR¹⁵;
- 40 X² is C(O), CR¹¹R¹¹, S, S(O) or S(O)₂;
- X³ is C(O), CR¹¹R¹¹, S(O)₂ or a bond;
- Y is X¹ or X²;
- Q is -CO₂R¹², -C(O)NHS(O)₂R¹⁴, -NHS(O)₂R¹⁴, -S(O)₂NHR¹⁵, -C(O)NR¹⁵R¹⁵, -CO₂R¹⁷, -C(O)NR¹⁸R²⁴, -CH₂OH, or 1H- or 2H-tetrazol-5-yl;
- 45 and the pharmaceutically acceptable salts thereof.
- A preferred embodiment of Formula I is that in which:
- R¹, R², R³ and R⁴ are hydrogen;
- R⁵ is X²-R⁷ or -OR⁶;
- R⁷ is R⁶;
- 50 R⁸ is R⁹;
- R¹⁰ is hydrogen or halogen;
- m is 0;
- n is 1 to 3;
- u is 0 in R⁶ and 1 in R⁹;
- v is 0;
- 55 X² is CR¹¹R¹¹ or S;
- Q is -CO₂R¹²; and the remaining substituents are as defined for Formula I;
- and the pharmaceutically acceptable salts thereof.

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Definitions

The following abbreviations have the indicated meanings:

- Me = methyl
- 5 Bz = benzyl
- Ph = phenyl
- t-Bu = tert-butyl
- i-Pr = isopropyl
- c-C₆H₁₁ = cyclohexyl
- 10 c-Pr = cyclopropyl
- c- = cyclo
- Ac = acetyl
- Tz = 5-tetrazolyl
- Th = 2- or 3- thienyl
- 15 c-C₅H₉ = cyclopentyl
- 1-Ad = 1-adamantyl.

Alkyl, alkenyl, and alkynyl are intended to include linear, branched, and cyclic structures and combinations thereof.

As used herein, the term "alkyl" includes "lower alkyl" and extends to cover carbon fragments having up to 20 carbon atoms. Examples of alkyl groups include octyl, nonyl, norbornyl, undecyl, dodecyl, tridecyl, tetradecyl, pentadecyl, eicosyl, 3,7-diethyl-2,2-dimethyl-4-propylnonyl, cyclododecyl, adamantyl, and the like.

As used herein, the term "lower alkyl" includes those alkyl groups of from 1 to 7 carbon atoms. Examples of lower alkyl groups include methyl, ethyl, propyl, isopropyl, butyl, sec- and tert-butyl, pentyl, 25 hexyl, heptyl, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, 2-methylcyclopropyl, cyclopropylmethyl, and the like.

The term "cycloalkyl" refers to a hydrocarbon ring having from 3 to 7 carbon atoms. Examples of cycloalkyl groups are cyclopropyl, cyclopentyl, cycloheptyl and the like.

"Lower alkenyl" groups include those alkenyl groups of 2 to 7 carbon atoms. Examples of lower alkenyl groups include vinyl, allyl, isopropenyl, pentenyl, hexenyl, heptenyl, cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclohexenyl, 1-propenyl, 2-butenyl, 2-methyl-2-butenyl and the like.

"Lower alkynyl" groups include those alkynyl groups of 2 to 7 carbon atoms. Examples of lower alkynyl groups include ethynyl, propargyl, 3-methyl-1-pentyne, 2-heptyne and the like.

As used herein, the term "lower alkoxy" includes those alkoxy groups of from 1 to 7 carbon atoms of a straight, branched, or cyclic configuration. Examples of lower alkoxy groups include methoxy, ethoxy, 35 propoxy, isopropoxy, cyclopropyloxy, cyclohexyloxy, and the like.

The term "monocyclic monoheterocyclic ring" which defines R²³ includes those monocyclic groups of 5 to 7 members containing only 1 heteroatom selected from N, S or O in the ring. Examples include tetrahydrofuran, tetrahydrothiophene, pyrrolidine, piperidine, tetrahydropyran, and the like.

40 The term "monocyclic or bicyclic heterocyclic ring" which defines R¹⁹ may be 2,5-dioxo-1-pyrrolidinyl, (3-pyridinylcarbonyl) amino, 1,3-dihydro-1,3-dioxo-2H-isoindol-2-yl, 1,3-dihydro-2H-isoindol-2-yl, 2,4-imidazolinedion-1-yl, 2,6-piperidinedion-1-yl, 2-imidazolyl, 2-oxo-1,3-dioxolen-4-yl, piperidin-1-yl, morpholin-1-yl, piperazin-1-yl and the like.

The point of attachment of any heterocyclic ring may be at any free valence of the ring.

45 The term standard amino acid is employed to include the following amino acids: alanine, asparagine, aspartic acid, arginine, cysteine, glutamic acid, glutamine, glycine, histidine, isoleucine, leucine, lysine, methionine, phenylalanine, proline, serine, threonine, tryptophan, tyrosine and valine. (See F.H.C. Crick, Symposium of the Society for Experimental Biology, 1958 (12) p. 140.)

It is understood that R¹ and R² may be located at any of positions 3,4,5,6,7 or 8 of the quinoline ring.

50 As used herein the term "lower alkylthio" includes those alkylthio groups of from 1 to 7 carbon atoms of a straight, branched or cyclic configuration. Examples of lower alkylthio groups include methylthio, propylthio, isopropylthio, cycloheptylthio, etc. By way of illustration, the propylthio group signifies -SCH₂CH₂CH₃.

The terms Ph(R¹⁰)₂ and Th(R¹⁰)₂ indicate a phenyl or thienyl group substituted with two R¹⁰ substituents.

Halogen includes F, Cl, Br, and I.

It is intended that the definitions of any substituent (e.g., R¹, R², R¹⁵, Ph(R¹⁰)₂, etc.) in a particular molecule be independent of its definitions elsewhere in the molecule. Thus, -NR¹⁵R¹⁵ represents -NHH,

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-NHCH₃, -NHC₆H₅, etc.

The monocyclic heterocyclic rings formed when two R¹⁵ groups join through N include pyrrolidine, piperidine, morpholine, thiamorpholine, piperazine, and N-methylpiperazine.

The prodrug esters of Q (i.e., when Q = CO₂R¹⁷) are intended to include the esters such as are described by Saari et al., J. Med. Chem., 21, No. 8, 746-753 (1978), Sakamoto et al., Chem. Pharm. Bull., 32, No. 6, 2241-2248 (1984) and Bundgaard et al., J. Med. Chem., 30, No. 3, 451-454 (1987).

Some of the compounds described herein contain one or more asymmetric centers and may thus give rise to diastereomers and optical isomers. The present invention is meant to comprehend such possible diastereomers as well as their racemic and resolved, enantiomerically pure forms and pharmaceutically acceptable salts thereof.

The pharmaceutical compositions of the present invention comprise a compound of Formula I as an active ingredient or a pharmaceutically acceptable salt, thereof, and may also contain a pharmaceutically acceptable carrier and optionally other therapeutic ingredients. The term "pharmaceutically acceptable salts" refers to salts prepared from pharmaceutically acceptable non-toxic bases including inorganic bases and organic bases. Salts derived from inorganic bases include aluminum, ammonium, calcium, copper, ferric, ferrous, lithium, magnesium, manganic salts, manganous, potassium, sodium, zinc and the like. Particularly preferred are the ammonium, calcium, magnesium, potassium and sodium salts. Salts derived from pharmaceutically acceptable organic non-toxic bases include salts of primary, secondary, and tertiary amines, substituted amines including naturally occurring substituted amines, cyclic amines and basic ion exchange resins, such as arginine, betaine, caffeine, choline, N,N'-dibenzylethylenediamine, diethylamine, 2-diethylaminoethanol, 2-dimethylaminoethanol, ethanolamine, ethylenediamine, N-ethylmorpholine, N-ethylpiperidine, glucamine, glucosamine, histidine, hydrabamine, isopropylamine, lysine, methylglucamine, morpholine, piperazine, piperidine, polyamine resins, procaine, purines, theobromine, triethylamine, trimethylamine, tripropylamine, tromethamine and the like.

When the compound of the present invention is basic, salts may be prepared from pharmaceutically acceptable non-toxic acids, including inorganic and organic acids. Such acids include acetic, benzenesulfonic, benzoic, camphorsulfonic, citric, ethanesulfonic, fumaric, gluconic, glutamic, hydrobromic, hydrochloric, isethionic, lactic, maleic, malic, mandelic, methanesulfonic, mucic, nitric, pamoic, pantothenic, phosphoric, succinic, sulfuric, tartaric, p-toluenesulfonic acid and the like. Particularly preferred are citric, hydrobromic, hydrochloric, maleic, phosphoric, sulfuric and tartaric acids.

It will be understood that in the discussion of methods of treatment which follows, references to the compounds of Formula I are meant to also include the pharmaceutically acceptable salts.

The ability of the compounds of Formula I to inhibit biosynthesis of the leukotrienes makes them useful for inhibiting the symptoms induced by the leukotrienes in a human subject. This inhibition of the mammalian biosynthesis of leukotrienes indicates that the compounds and pharmaceutical compositions thereof are useful to treat, prevent, or ameliorate in mammals and especially in humans: 1) pulmonary conditions including diseases such as asthma, 2) allergies and allergic reactions such as allergic rhinitis, contact dermatitis, allergic conjunctivitis, and the like, 3) inflammation such as arthritis or inflammatory bowel disease, 4) pain, 5) skin conditions such as psoriasis and the like, 6) cardiovascular conditions such as angina, endotoxin shock, and the like and 7) renal insufficiency arising from ischaemia induced by immunological or chemical (cyclosporin) etiology, and that the compounds are cytoprotective agents.

The cytoprotective activity of a compound may be observed in both animals and man by noting the increased resistance of the gastrointestinal mucosa to the noxious effects of strong irritants, for example, the ulcerogenic effects of aspirin or indomethacin. In addition to lessening the effect of non-steroidal anti-inflammatory drugs on the gastrointestinal tract, animal studies show that cytoprotective compounds will prevent gastric lesions induced by oral administration of strong acids, strong bases, ethanol, hypertonic saline solutions and the like.

Two assays can be used to measure cytoprotective ability. These assays are; (A) an ethanol-induced lesion assay and (B) an indomethacin-induced ulcer assay and are described in EP 140,684.

The magnitude of prophylactic or therapeutic dose of a compound of Formula I will, of course, vary with the nature of the severity of the condition to be treated and with the particular compound of Formula I and its route of administration. It will also vary according to the age, weight and response of the individual patient. In general, the daily dose range for anti-asthmatic, anti-allergic or anti-inflammatory use and generally, uses other than cytoprotection, lie within the range of from about 0.001 mg to about 100 mg per kg body weight of a mammal, preferably 0.01 mg to about 10 mg per kg, and most preferably 0.1 to 1 mg per kg, in single or divided doses. On the other hand, it may be necessary to use dosages outside these limits in some cases.

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For use where a composition for intravenous administration is employed, a suitable dosage range for anti-asthmatic, anti-inflammatory or anti-allergic use is from about 0.001 mg to about 25 mg (preferably from 0.01 mg to about 1 mg) of a compound of Formula I per kg of body weight per day and for cytoprotective use from about 0.1 mg to about 100 mg (preferably from about 1 mg to about 100 mg and more preferably from about 1 mg to about 10 mg) of a compound of Formula I per kg of body weight per day.

In the case where an oral composition is employed, a suitable dosage range for anti-asthmatic, anti-inflammatory or anti-allergic use is, e.g. from about 0.01 mg to about 100 mg of a compound of Formula I per kg of body weight per day, preferably from about 0.1 mg to about 10 mg per kg and for cytoprotective use from 0.1 mg to about 100 mg (preferably from about 1 mg to about 100 mg and more preferably from about 10 mg to about 100 mg) of a compound of Formula I per kg of body weight per day.

For the treatment of diseases of the eye, ophthalmic preparations for ocular administration comprising 0.001-1% by weight solutions or suspensions of the compounds of Formula I in an acceptable ophthalmic formulation may be used.

The exact amount of a compound of the Formula I to be used as a cytoprotective agent will depend on, inter alia, whether it is being administered to heal damaged cells or to avoid future damage, on the nature of the damaged cells (e.g., gastrointestinal ulcerations vs. nephrotic necrosis), and on the nature of the causative agent. An example of the use of a compound of the Formula I in avoiding future damage would be co-administration of a compound of the Formula I with a non-steroidal anti-inflammatory drug (NSAID) that might otherwise cause such damage (for example, indomethacin). For such use, the compound of Formula I is administered from 30 minutes prior up to 30 minutes after administration of the NSAID. Preferably it is administered prior to or simultaneously with the NSAID, (for example, in a combination dosage form).

Any suitable route of administration may be employed for providing a mammal, especially a human with an effective dosage of a compound of the present invention. For example, oral, rectal, topical, parenteral, ocular, pulmonary, nasal, and the like may be employed. Dosage forms include tablets, troches, dispersions, suspensions, solutions, capsules, creams, ointments, aerosols, and the like.

The pharmaceutical compositions of the present invention comprise a compound of formula I as an active ingredient or a pharmaceutically acceptable salt thereof, and may also contain a pharmaceutically acceptable carrier and optionally other therapeutic ingredients. The term "pharmaceutically acceptable salts" refers to salts prepared from pharmaceutically acceptable non-toxic bases or acids including inorganic bases or acids and organic bases or acids.

The compositions include compositions suitable for oral, rectal, topical, parenteral (including subcutaneous, intramuscular, and intravenous), ocular (ophthalmic), pulmonary (nasal or buccal inhalation), or nasal administration, although the most suitable route in any given case will depend on the nature and severity of the conditions being treated and on the nature of the active ingredient. They may be conveniently presented in unit dosage form and prepared by any of the methods well-known in the art of pharmacy.

For administration by inhalation, the compounds of the present invention are conveniently delivered in the form of an aerosol spray presentation from pressurized packs or nebulisers. The compounds may also be delivered as powders which may be formulated and the powder composition may be inhaled with the aid of an insufflation powder inhaler device. The preferred delivery system for inhalation is a metered dose inhalation (MDI) aerosol, which may be formulated as a suspension or solution of compound I in suitable propellants, such as fluorocarbons or hydrocarbons.

Suitable topical formulations of compound I include transdermal devices, aerosols, creams, ointments, lotions, dusting powders, and the like.

In practical use, the compounds of formula I can be combined as the active ingredient in intimate admixture with a pharmaceutical carrier according to conventional pharmaceutical compounding techniques. The carrier may take a wide variety of forms depending on the form of preparation desired for administration, e.g., oral or parenteral (including intravenous). In preparing the compositions for oral dosage form, any of the usual pharmaceutical media may be employed, such as, for example, water, glycols, oils, alcohols, flavoring agents, preservatives, coloring agents and the like in the case of oral liquid preparations, such as, for example, suspensions, elixirs and solutions; or carriers such as starches, sugars, microcrystalline cellulose, diluents, granulating agents, lubricants, binders, disintegrating agents and the like in the case of oral solid preparations such as, for example, powders, capsules and tablets, with the solid oral preparations being preferred over the liquid preparations. Because of their ease of administration, tablets and capsules represent the most advantageous oral dosage unit form in which case solid pharmaceutical carriers are obviously employed. If desired, tablets may be coated by standard aqueous or nonaqueous techniques.

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In addition to the common dosage forms set out above, the compounds of formula I may also be administered by controlled release means and/or delivery devices such as those described in U.S. Patent Nos. 3,845,770; 3,916,899; 3,536,809; 3,598,123; 3,630,200 and 4,008,719.

Pharmaceutical compositions of the present invention suitable for oral administration may be presented as discrete units such as capsules, cachets or tablets each containing a predetermined amount of the active ingredient, as a powder or granules or as a solution or a suspension in an aqueous liquid, a non-aqueous liquid, an oil-in-water emulsion or a water-in-oil liquid emulsion. Such compositions may be prepared by any of the methods of pharmacy but all methods include the step of bringing into association the active ingredient with the carrier which constitutes one or more necessary ingredients. In general, the compositions are prepared by uniformly and intimately admixing the active ingredient with liquid carriers or finely divided solid carriers or both, and then, if necessary, shaping the product into the desired presentation. For example, a tablet may be prepared by compression or molding, optionally with one or more accessory ingredients. Compressed tablets may be prepared by compressing in a suitable machine, the active ingredient in a free-flowing form such as powder or granules, optionally mixed with a binder, lubricant, inert diluent, surface active or dispersing agent. Molded tablets may be made by molding in a suitable machine, a mixture of the powdered compound moistened with an inert liquid diluent. Desirably, each tablet contains from about 2.5 mg to about 500 mg of the active ingredient and each cachet or capsule contains from about 2.5 to about 500 mg of the active ingredient.

The following are examples of representative pharmaceutical dosage forms for the compounds of Formula I:

Injectable Suspension (I.M.)	mg/ml
Compound of Formula I	10
Methylcellulose	5.0
Tween 80	0.5
Benzyl alcohol	9.0
Benzalkonium chloride	1.0
Water for injection to a total volume of 1 ml	

Tablet	mg/tablet
Compound of Formula I	25
Microcrystalline Cellulose	415
Providone	14.0
Pregelatinized Starch	43.5
Magnesium Stearate	2.5
	500

Capsule	mg/capsule
Compound of Formula I	25
Lactose Powder	573.5
Magnesium Stearate	1.5
	600

Aerosol	Per canister
Compound of Formula I	24 mg
Lecithin, NF Liquid Concentrate	1.2 mg
Trichlorofluoromethane, NF	4.025 gm
Dichlorodifluoromethane, NF	12.15 gm

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In addition to the compounds of Formula I, the pharmaceutical compositions of the present invention can also contain other active ingredients, such as cyclooxygenase inhibitors, non-steroidal anti-inflammatory drugs (NSAIDs), peripheral analgesic agents such as zomepirac diflunisal and the like. The weight ratio of the compound of the Formula I to the second active ingredient may be varied and will depend upon the effective dose of each ingredient. Generally, an effective dose of each will be used. Thus, for example, when a compound of the Formula I is combined with an NSAID the weight ratio of the compound of the formula I to the NSAID will generally range from about 1000:1 to about 1:1000, preferably about 200:1 to about 1:200. Combinations of a compound of the Formula I and other active ingredients will generally also be within the aforementioned range, but in each case, an effective dose of each active ingredient should be used.

NSAIDs can be characterized into five groups:

- (1) the propionic acid derivatives;
 - (2) the acetic acid derivatives;
 - (3) the fenamic acid derivatives;
 - (4) the biphenylcarboxylic acid derivatives; and
 - (5) the oxicams
- or a pharmaceutically acceptable salt thereof.

The propionic acid derivatives which may be used comprise: alminoprofen, benoxaprofen, bucloxic acid, carprofen, fenbufen, fenoprofen, fluprofen, flurbiprofen, ibuprofen, indoprofen, ketoprofen, miroprofen, naproxen, oxaprozin, piroprofen, prano-profen, suprofen, tiaprofenic acid, and tioprofen. Structurally related propionic acid derivatives having similar analgesic and anti-inflammatory properties are also intended to be included in this group.

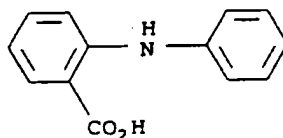
Thus, "propionic acid derivatives" as defined herein are non-narcotic analgesics/non-steroidal anti-inflammatory drugs having a free $-\text{CH}(\text{CH}_3)\text{COOH}$ or $-\text{CH}_2\text{CH}_2\text{COOH}$ group (which optionally can be in the form of a pharmaceutically acceptable salt group, e.g., $-\text{CH}(\text{CH}_3)\text{COO}^-\text{Na}^+$ or $-\text{CH}_2\text{CH}_2\text{COO}^-\text{Na}^+$), typically attached directly or via a carbonyl function to a ring system, preferably to an aromatic ring system.

The acetic acid derivatives which may be used comprise: indomethacin, which is a preferred NSAID, acetaminophen, alclufenac, clidanac, diclofenac, fenclofenac, fenclozic acid, fentiazac, furofenac, ibufenac, isoxepac, oxpinac, sulindac, tiopinac, tolmetin, zidometacin and zomepirac. Structurally related acetic acid derivatives having similar analgesic and anti-inflammatory properties are also intended to be encompassed by this group.

Thus, "acetic acid derivatives" as defined herein are non-narcotic analgesics/non-steroidal anti-inflammatory drugs having a free $-\text{CH}_2\text{COOH}$ group (which optionally can be in the form of a pharmaceutically acceptable salt group, e.g. $-\text{CH}_2\text{COO}^-\text{Na}^+$), typically attached directly to a ring system, preferably to an aromatic or heteroaromatic ring system.

The fenamic acid derivatives which may be used comprise: flufenamic acid, meclofenamic acid, mefenamic acid, niflumic acid and tolfenamic acid. Structurally related fenamic acid derivatives having similar analgesic and anti-inflammatory properties are also intended to be encompassed by this group.

Thus, "fenamic acid derivatives" as defined herein are non-narcotic analgesics/non-steroidal anti-inflammatory drugs which contain the basic structure:

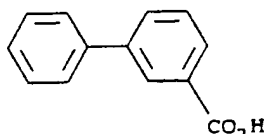


which can bear a variety of substituents and in which the free $-\text{COOH}$ group can be in the form of a pharmaceutically acceptable salt group, e.g., $-\text{COO}^-\text{Na}^+$.

The biphenylcarboxylic acid derivatives which can be used comprise: diflunisal and flufenisal. Structurally related biphenylcarboxylic acid derivatives having similar analgesic and anti-inflammatory properties are also intended to be encompassed by this group.

Thus, "biphenylcarboxylic acid derivatives" as defined herein are non-narcotic analgesics/non-steroidal anti-inflammatory drugs which contain the basic structure:

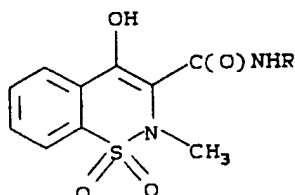
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which can bear a variety of substituents and in which the free -COOH group can be in the form of a pharmaceutically acceptable salt group, e.g., -COO⁻Na⁺.

The oxicams which can be used in the present invention comprise: isoxicam, piroxicam, sudoxicam and tenoxicam. Structurally related oxicams having similar analgesic and anti-inflammatory properties are also intended to be encompassed by this group.

Thus, "oxicams" as defined herein are non narcotic analgesics/non-steroidal anti-inflammatory drugs which have the general formula:



wherein R is an aryl or heteroaryl ring system.

The following NSAIDs may also be used: amfenac sodium, aminoprofen, anitrazafen, antrafenine, auranofin, bendazac lysinate, benzydanine, beprozoin, broperamol, bufexamac, cinmetacin, ciproquazone, cloximate, dazidamine, deboxamet, delmetacin, detomidine, dexindoprofen, diacerein, di-fisalamine, difenpyramide, emorfazone, enfenamic acid, enolicam, eprizole, etersalate, etodolac, etofenamate, fanetizole mesylate, fenclorac, fendosal, fenflumizole, feprazone, floctafenine, flunixin, flunoxaprofen, fluproquazone, fopirtoline, fosfosal, furciprofen, glucametacin, guaimesal, ibuprofen, isofezolac, isonixim, isoprofen, isoxicam, lefetamine HCl, leflunomide, lofenizole, lonazolac calcium, lotifazole, loxoprofen, lysin clonixinate, meclofenamate sodium, meseclazone, nabumetone, nictindole, nimesulide, orpanoxin, oxametacin, oxapadol, perisoxal citrate, pimeprofen, pimetacin, piroxifen, pirazolac, pirfenidone, proglumetacin maleate, proquazone, pyridoxiprofen, sudoxicam, talmecacin, talniflumate, tenoxicam, thiazolinobutazone, thielavin B, tiaramide HCl, tiffamizole, timegadine, tolpadol, tryptamid and ufenamate.

The following NSAIDs, designated by company code number (see e.g., Pharmaprojects), may also be used:

480156S, AA861, AD1590, AFP802, AFP860, AI77B, AP504, AU8001, BPPC, BW540C, CHINOIN 127, CN100, EB382, EL508, F1044, GV3658, ITF182, KCNTEI6090, KME4, LA2851, MR714, MR897, MY309, ONO3144, PR823, PV102, PV108, R830, RS2131, SCR152, SH440, SIR133, SPAS510, SQ27239, ST281, SY6001, TA60, TAI-901 (4-benzoyl-1-indancarboxylic acid), TVX2706, U60257, UR2301, and WY41770.

Finally, NSAIDs which may also be used include the salicylates, specifically acetyl salicylic acid and the phenylbutazones, and pharmaceutically acceptable salts thereof.

In addition to indomethacin, other preferred NSAIDs are acetyl salicylic acid, diclofenac, fenbufen, fenoprofen, flurbiprofen, ibuprofen, ketoprofen, naproxen, phenylbutazone, piroxicam, sulindac and tolmetin.

Pharmaceutical compositions comprising the Formula I compounds may also contain inhibitors of the biosynthesis of the leukotrienes such as are disclosed in EP 138,481 (April 24, 1985), EP 115,394 (August 8, 1984), EP 136,893 (April 10, 1985), and EP 140,709 (May 8, 1985).

The compounds of the formula I may also be used in combination with leukotriene antagonists such as those disclosed in EP 106,565 (April 25, 1984) and EP 104,885 (April 4, 1984) which are hereby incorporated herein by reference and others known in the art such as those disclosed in EP Application Nos. 56,172 (July 21, 1982) and 61,800 (June 10, 1982); and in U.K. Patent Specification No. 2,058,785 (April 15, 1981).

Pharmaceutical compositions comprising the Formula I compounds may also contain as the second active ingredient, prostaglandin antagonists such as those disclosed in EP 11,067 (May 28, 1980) or thromboxane antagonists such as those disclosed in U.S. Pat. 4,237,160. They may also contain histidine

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decarboxylase inhibitors such as α -fluoromethylhistidine, described in U.S. Pat. 4,325,961. The compounds of the formula I may also be advantageously combined with an H₁ or H₂-receptor antagonist, such as for instance acetamazole, aminothiadiazoles disclosed in EP 40,696 (December 2, 1981), benadryl, cimetidine, famotidine, framamine, histadyl, phenergan, ranitidine, terfenadine and like compounds, such as those
 5 disclosed in U.S. Patent Nos. 4,283,408; 4,362,736; and 4,394,508. The pharmaceutical compositions may also contain a K⁺/H⁺ ATPase inhibitor such as omeprazole, disclosed in U.S. Pat. 4,255,431, and the like. Compounds of Formula I may also be usefully combined with most cell stabilizing agents, such as 1,3-bis-(2-carboxychromon-5-yloxy)-2-hydroxypropane and related compounds described in British Patent Specifications 1,144,905 and 1,144,906. Another useful pharmaceutical composition comprises the Formula I
 10 compounds in combination with serotonin antagonists such as methysergide, the serotonin antagonists described in Nature, Vol. 316, pages 126-131, 1985, and the like. Each of the references referred to in this paragraph is hereby incorporated herein by reference.

Other advantageous pharmaceutical compositions comprise the Formula I compounds in combination with anti-cholinergics such as ipratropium bromide, bronchodilators such as the beta agonist salbutamol,
 15 metaproterenol, terbutaline, fenoterol and the like, and the anti-asthmatic drugs theophylline, choline theophyllinate and enprofylline, the calcium antagonists nifedipine, diltiazem, nitrendipine, verapamil, nimodipine, felodipine, etc. and the corticosteroids, hydrocortisone, methylprednisolone, betamethasone, dexamethasone, beclomethasone, and the like.

Compounds of the present invention can be prepared according to the following methods. Temperatures are in degree Celsius.

The starting methoxy phenylhydrazines II are either commercially available or are described in the chemical literature as are the acetamidophenols XXVI. The benzyl phenylhydrazine starting materials III are prepared as described in EP 166,591 (17102 IA) and the ketones IV and XXXI are prepared as described in EP 166,591 and EP 275,667 (17496 IA). The 2-(halomethyl)quinolines VII are available from literature
 25 methods described in "Quinolines" Parts I and II, G. Jones (ED.), John Wiley & Sons, Toronto, 1977 and 1982. The preparation of VII by halogenation of the corresponding 2-methylquinolines is also described in the Jones' volumes. The benzyl halides, (R¹⁰)₂ PhCH₂-Hal, are readily prepared and many such compounds are described in the prior art, such as U.S. Patent 4,808,608 (17323 IB). Hal in VII and (R¹⁰)₂ PhCH₂-Hal represents Cl, Br or I.

Many syntheses of indoles are well-known in the chemical literature: see for example, "Heterocyclic compounds" Volume 25, Parts I, II, III, W.J. Houlihan (Ed.), Interscience, J. Wiley & Sons, N.Y., 1979, and "The Chemistry of Indoles" by R.J. Sundberg, Academic Press, N.Y., 1970. One of the most common syntheses is known as the Fischer Indole Synthesis, and is abbreviated in the following methods as "Fischer".

The -CO₂H and -CO₂R¹² groups in the intermediates and final products in the various methods can be transformed to other representatives of Q such as -CONHS(O)₂R¹⁴, -NHS(O)₂R¹⁴, -CONR¹⁵R¹⁵, -CH₂OH or tetrazol-5-yl by the methodology described in U.S. Patent 4,808,608 (17323IV). The preparation of the pro-drug forms (Q is -CO₂R¹⁷) from the acids may be effected by the methodology of EP 104,885 (16830 IA).

It will be apparent to one skilled in the art that the various functional groups (R¹, R², Y, Q, etc.) must be
 40 chosen so as to be compatible with the chemistry being carried out. Such compatibility can often be achieved by protecting groups, or by specific variations in the sequence of the reactions.

When R⁵ is S-R⁷, the corresponding sulfoxides and sulfones can be prepared by oxidation of the sulfides with one or two equivalents of an oxidizing agent such as m-chloroperbenzoic acid or monoperox-yphthalic acid or oxone (Trost, J. Org. Chem., 1988, pg.532).

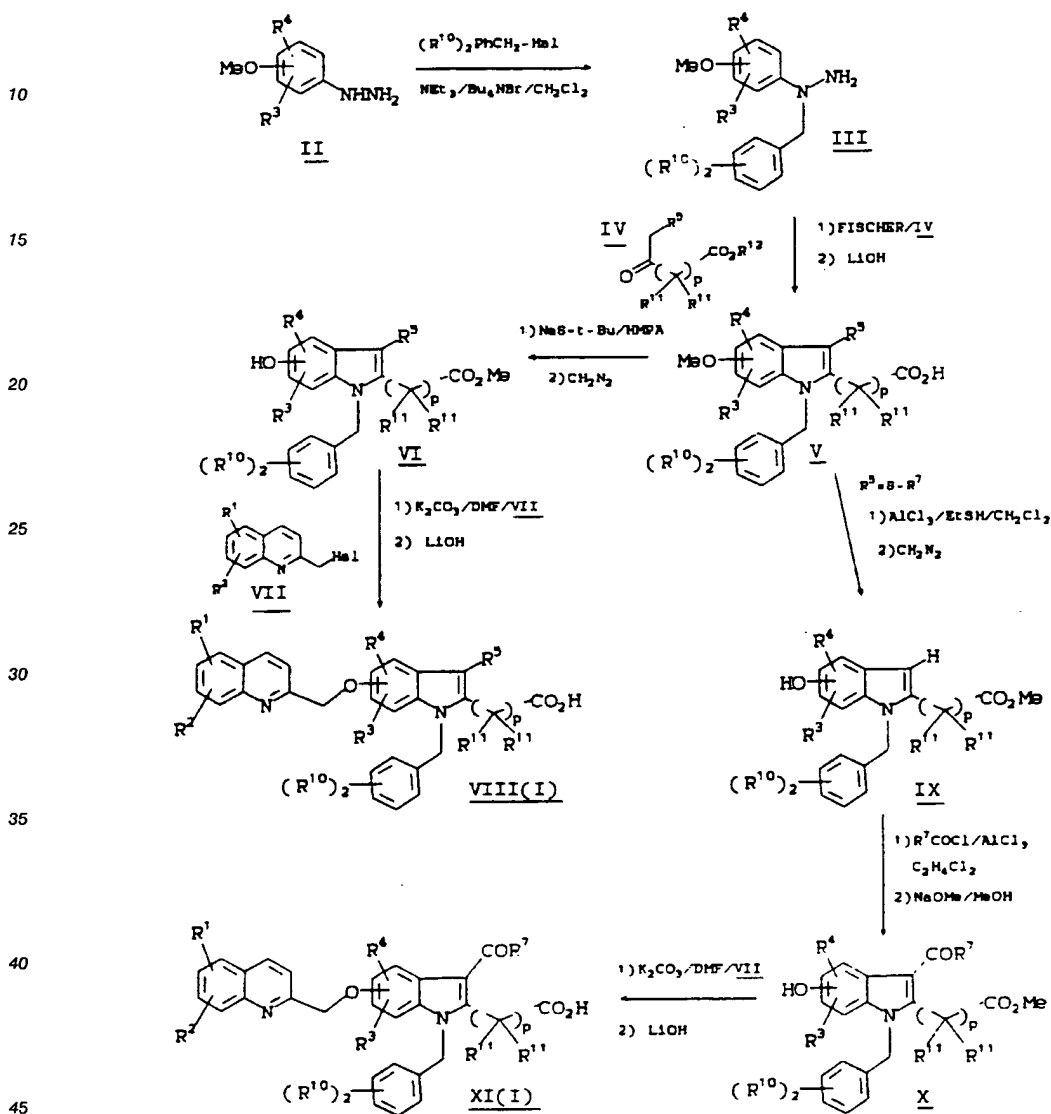
Many of the following methods involve a basic hydrolysis of an ester function to obtain the corresponding carboxylic acid. In all cases, the free acid is obtained by acidification of the reaction mixture with a suitable acid such as hydrochloric, sulfuric, acetic, trifluoroacetic acid, etc.

Compounds VIII, XI, XV, XIX, XXXVI and their precursor esters are all examples of the Formula I compounds of the present invention.

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Method 1

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Method 1

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Intermediate V is prepared by a Fischer reaction between benzylphenylhydrazine III and ketone IV, followed by hydrolysis with an aqueous solution of an alkali hydroxide or other suitable hydroxide in mixture with a suitable water miscible organic solvent such as tetrahydrofuran (THF) or methanol (MeOH). The methoxy acid V is demethylated by heating with an alkali salt of an aliphatic thiol in a suitable solvent such as hexamethylphosphorotriamide (HMPA) or N-methylpyrrolidone (NMP). The reaction mixture is acidified and the crude acid so obtained is converted to the methyl ester VI by treatment with diazomethane. The phenol VI is coupled to the 2-halomethylquinoline VII, by stirring with a base (preferably an alkali hydride or carbonate) in a suitable solvent such as dimethyl formamide (DMF), NMP, acetone or the like. The resulting

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ester is hydrolysed by base to yield VIII, a compound of the present invention.

When intermediate V contains a sulfide group attached to position 3, treatment with a Lewis acid, such as $AlCl_3$, and an aliphatic thiol, simultaneously effects demethylation and removes the sulfide group. Suitable solvents for this reaction are methylene chloride, 1,2-dichloroethane, etc. The resulting acid is then converted to the methyl ester IX with diazomethane. A Friedel-Crafts reaction between IX and an acid chloride, R^7COCl , simultaneously introduces the acyl substituent into the 3-position of the indole ring and onto the phenolic hydroxyl group. The acyl group is removed from the phenol by treatment with sodium methoxide in MeOH to yield acylphenol X. Phenol X is coupled with VII as described for the coupling of VI and VII above. In these coupling reactions, it is at times advantageous to add a catalyst such as potassium iodide or tetraethylammonium bromide, especially when Hal is chlorine. A final hydrolysis yields compound XI.

METHOD 2

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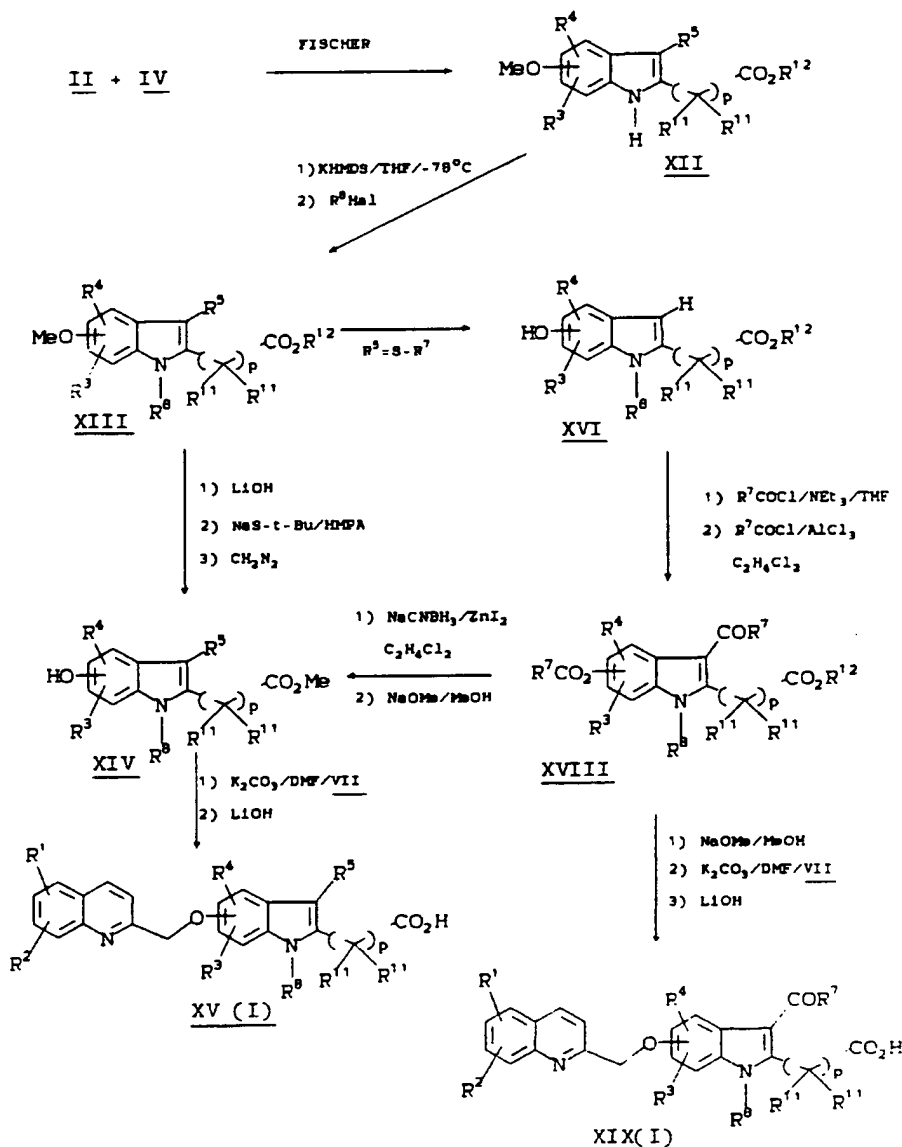
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Method 2

Intermediate XII is prepared by a Fischer reaction between methoxyphenyl hydrazine II and ketone IV, followed by alkylation of the indole nitrogen, after deprotonation using potassium hexamethyldisilazane in an ether solvent such as tetrahydrofuran (THF), with an alkyl or aralkyl halide.

The methoxy group in XIII is removed using the conditions of Method 1. The corresponding phenol XIV is now coupled with the 2-halomethylquinoline VII by stirring with a base (preferably an alkali hydride or carbonate) in a suitable solvent such as DMF, NMP or the like. The resulting ester is hydrolysed using base to yield XV a compound of the present invention.

When intermediate XIII contains a sulfide at position 3, treatment with a Lewis acid such as AlCl_3 and an aliphatic thiol simultaneously effects demethylation and removes the sulfide group. Suitable solvents for this reaction are dichloromethane or dichloroethane. In a variation of Method 1, the phenolic hydroxyl in XVI is first acylated with the reagent R^7COCl (XVII) in the presence of a weak base such as triethylamine. A Friedel-Crafts reaction is then carried out on the O-acylated intermediate, with an additional mole of XVII and AlCl_3 , to yield the intermediate XVIII. Acyl ester XVIII may then be reduced to a 3-alkyl indole XIV using sodium cyanoborohydride in dichloroethane using a zinc iodide catalyst.

Acyl ester XVIII is cleaved to the indole phenol by hydrolysis with sodium methoxide in methanol and is coupled to 2-halomethyl quinoline VII using a base such as an alkali hydride or carbonate in a solvent such as DMF or NMP. Hydrolysis of the resulting compound using base yields the compound of the present invention XIX.

METHOD 3

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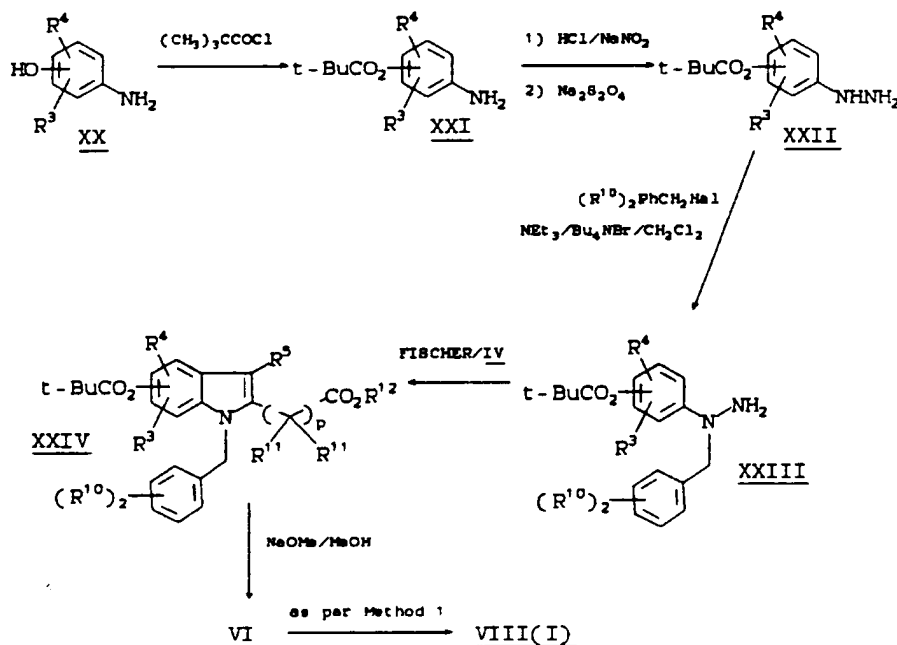
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Method 3

A suitably substituted aminophenol XX is protected on oxygen by the use of pivaloyl chloride dissolved in CH_2Cl_2 using triethyl amine as base. The pivaloate ester XXI is then diazotized using hydrochloric acid and sodium nitrite in an aqueous solvent and the transient diazonium species reduced in situ to the hydrazine XXII using sodium hydrosulfite in water. Benzylation of the hydrazine is effected as described in Method 1.

The O-pivaloyl-N-benzylhydrazine XXIII is subjected to a Fischer indolization using the appropriate ketone IV to produce the indole XXIV. Cleavage of the O-pivaloyl group using sodium methoxide in methanol transforms the product into the phenolic indole VI which is converted to the products of this invention as described in Method 1.

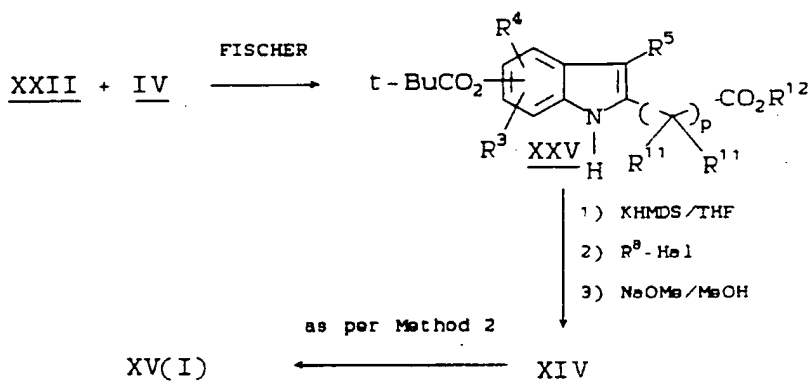
METHOD 4

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Method 4

The pivaloyloxyphenylhydrazine XXII is used directly in the Fischer indolization using ketone IV. N-Alkylation of the indole XXV, as described in Method 2, followed by removal of the pivaloyl group as described, yields the phenolic indole XIV which is converted as described in Method 2 to the products of this invention.

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Method 5

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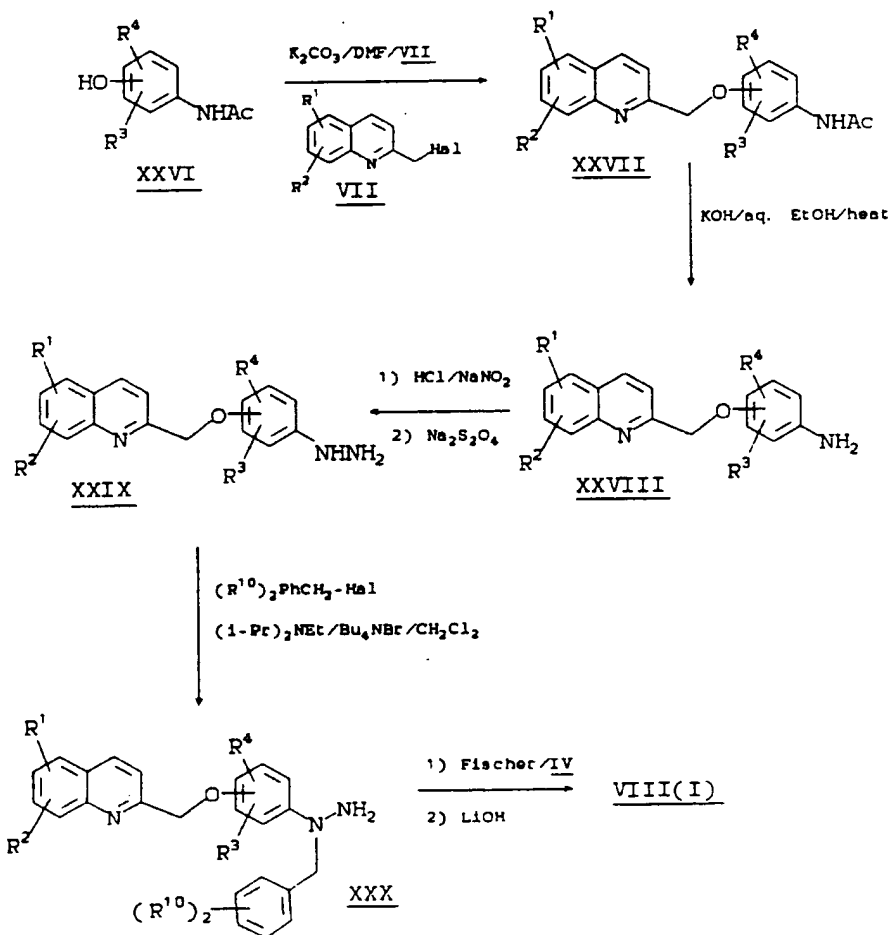
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Method 5

A suitable N-acetylated aminophenol XXVI is reacted with VII using an alkali hydride or carbonate, such as potassium carbonate as a base in a polar solvent like DMF or NMP. The quinolinylmethoxy acetanilide XXVII is then de-acetylated using standard basic conditions, preferably using alcoholic potassium hydroxide under reflux to produce the quinolinylmethoxy aniline derivative XXVIII. Conversion of the quinolinylmethoxy aniline derivative to the hydrazine analogue XXIX is effected through reduction of the intermediate diazonium salt using sodium hydrosulfite in an aqueous medium.

The hydrazine XXIX is then N-benzylated using a benzyl halide in an organic solvent such as methylene chloride containing an amine base such as diisopropylethylamine and preferably tetra-n-butylammonium bromide as catalyst.

The hydrazine XXX is then processed using a Fischer indolization with ketone IV according to Methods 1, 2, 3 and 4 to produce compounds of the present invention.

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Method 6

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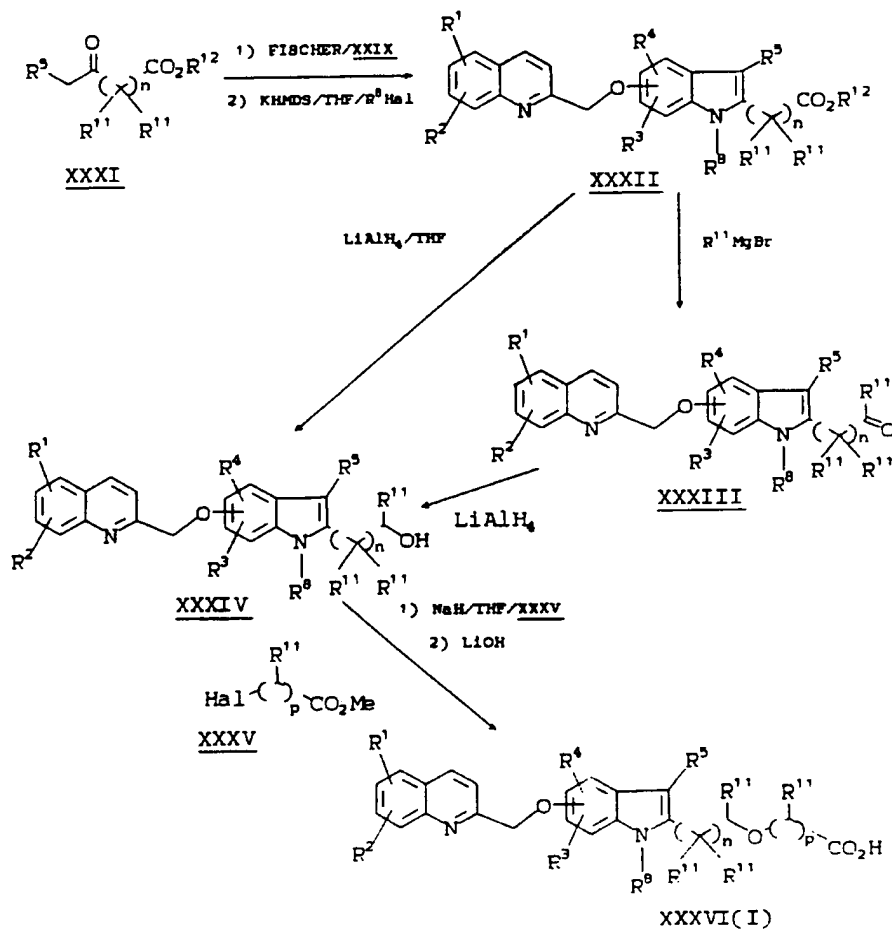
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Method 6

Hydrazine XXIX may also be transformed directly to unsubstituted indoles by a Fischer reaction with various ketones like XXXI. N-Alkylation of the indoles is effected using the conditions described in Method 2 to produce quinolinylmethoxyindole alkanoate esters XXXII. Such esters are transformed to ketones or carbinols via Grignard conditions using alkyl magnesium halides in ether solvents like diethyl ether or through the use of lithium aluminum hydride in ether solvents like THF. The carbinols XXXIV so produced may be further transformed into ester compounds of the present invention by reacting with α -halo esters XXXV using sodium hydride as base in a suitable solvent like THF. Subsequent hydrolysis of the esters using Method 1 leads to acid compounds of the present invention.

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Method 7

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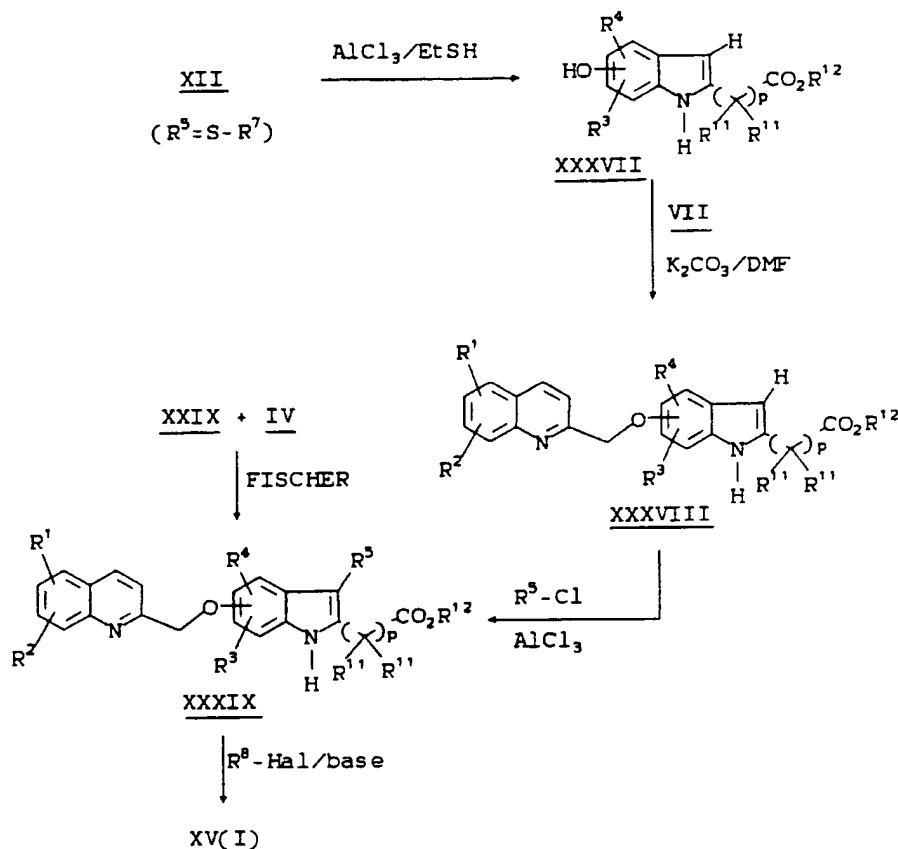
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METHOD 7

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Phenol XXXVII is obtained by treatment of XII ($\text{R}^5 = \text{S}-\text{R}^7$) with a Lewis acid and a thiol, as in Method 1 for the conversion of V to IX. Compound XXXVIII is then obtained by reaction of XXXVII with VII in the presence of a base in a suitable solvent, as described for the conversion of VI to VIII in Method 1. The introduction of R^5 in XXXIX is conveniently effected by an electrophilic reaction between XXXVIII and R^5-Cl (R^5 not = X^1-R^6). Such reactions are frequently catalysed by Lewis acids or proton acids such as AlCl_3 , SnCl_4 , TiCl_4 , BBr_3 , HCl , HBr and the like. They may be carried out in a variety of solvents, with a preference for non-protonic solvents such as dichloromethane, 1,2-dichloroethane, nitromethane, chlorobenzene and the like. It will be obvious to one skilled in the art, that the chlorine in R^5-Cl , in this and the other Methods, may often be replaced by another halogen or by a hydroxyl group, or R^5-Cl may be replaced by an acid anhydride $(\text{R}^7\text{CO})_2\text{O}$. An alternative synthesis of XXXIX is to effect a Fischer reaction between compounds IV and XXIX. Introduction of R^8 into XXXIX, is accomplished by alkylation with R^8-Hal and a base as described previously for Methods 2, 4 and 6. Finally, hydrolysis of the ester will yield XV. Alternatively, the ester group in XXXIX can be hydrolysed, and the corresponding free acid ($\text{R}^{12} = \text{H}$) alkylated on the indole nitrogen with R^8-Hal and an aqueous base, such as NaOH , and a phase-transfer catalyst, such as

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methyltriocetylammmonium chloride. Alkylation of the acid corresponding to XXXIX ($R^{12} = H$) can also be effected using a strong base such as sodium hydride in a solvent such as DMF. This latter procedure usually gives the ester of XV in which the carboxyl group has also been alkylated. The free acid XV can be obtained by standard hydrolysis procedures. If R^8 in XV or the ester precursor of XV is alkenyl, it can be reduced to alkyl using hydrogen gas, and a Pt or Pd catalyst in a suitable solvent, at atmospheric pressure.

Method 8

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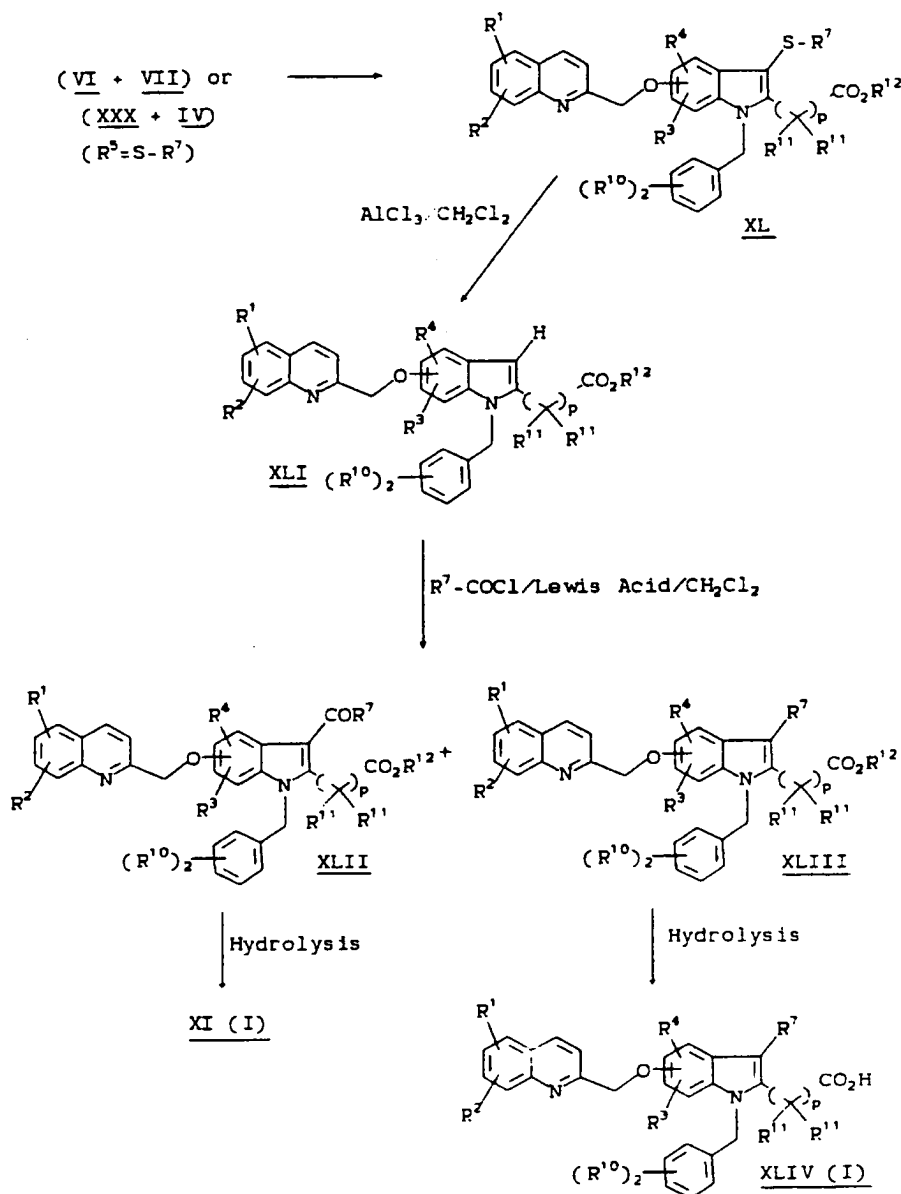
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METHOD 8

Compound XL may be prepared either by the coupling of VI to VII (Method 1) or by a Fischer reaction between IV and XXX (Method 5). Compound XL may be desulfurized by treatment with a Lewis acid such as AlCl_3 , or by reduction with Raney nickel, to give compound XLI. A Friedel-Crafts reaction on XLI with the reagent R^7COCl and a Lewis acid catalyst such as AlCl_3 yields the 3-acyl derivative XLII, hydrolysis of which yields XI. In the Friedel-Crafts reaction, carbon monoxide may be lost and compound XLIII is formed; hydrolysis under standard conditions then yields XLIV. The formation of XLIII occurs when the cation R^{7+} is especially stable and when the reagents R^7COCl and the Lewis acid are mixed before adding XLI. If the Lewis acid is added last, the main product is usually the acylated compound XLII. If a milder Lewis acid such as TiCl_4 is used, the main product is also XLII.

It will be obvious to one skilled in the art that the reagent R^7COCl can often be replaced by $\text{R}^7\text{CO-Hal}$ (Hal = F, Br or I) or $(\text{R}^7\text{CO})_2\text{O}$.

Representative Compounds

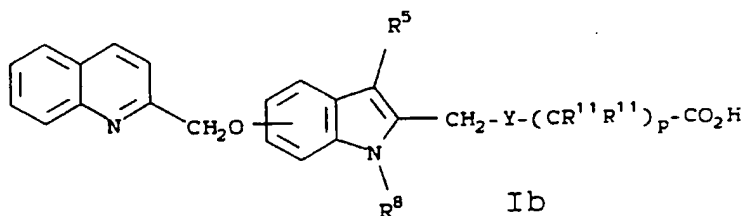
Table I and Table II illustrate compounds having the formulae Ia and Ib respectively representative of the present invention. "Attach point" is the position on the indole nucleus where the quinolylmethoxy moiety is attached.

TABLE I

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TABLE II



Ex No.	ATTACH POINT	R ⁸	R ⁵	Y-(CR ¹¹ R ¹¹) _p
19	6	-CH ₂ Ph-4-Cl	-S-t-Bu	C(Me) ₂
20	7	-CH ₂ Ph-4-Cl	-S-t-Bu	C(Me) ₂
22	5	-CH ₂ Ph-4-Cl	-S-t-Bu	CH ₂ OCH(Me)
23	4	-CH ₂ Ph-4-Cl	H	C(Me) ₂
24	6	Me	-C(O)Ph-4-Cl	C(Me) ₂
25	6	Me	-CH ₂ Ph-4-Cl	C(Me) ₂
26	5	-CH ₂ Ph-4-Cl	-O-i-Pr	C(Me) ₂
27	5	-CH ₂ Ph-4-Cl	-S-t-Bu	CH(Et)
28	5	-CH ₂ Ph-4-Cl	-C(O)-CF ₃	C(Me) ₂
29	5	-CH ₂ Ph-4-Cl	-C(O)CH ₂ -t-Bu	CH(Me)
30	5	H	-C(O)CH ₂ -t-Bu	C(Me) ₂
31	5	-CH ₂ Ph-4-CF ₃	-C(O)CH ₂ -t-Bu	C(Me) ₂
32	5	-CH ₂ Ph	-C(O)CH ₂ -t-Bu	C(Me) ₂
33	5	-CH ₂ Ph-3-OMe	-C(O)CH ₂ -t-Bu	C(Me) ₂
34	5	-CH ₂ CHCH ₂	-C(O)CH ₂ -t-Bu	C(Me) ₂
35	5	-CH ₂ Ph-4-OMe	-C(O)CH ₂ -t-Bu	C(Me) ₂
36	5	Me	-C(O)CH ₂ -t-Bu	C(Me) ₂
37	6	H	-CH ₂ Ph-4-Cl	C(Me) ₂
38	6	-S(O) ₂ Ph	-CH ₂ Ph-4-Cl	C(Me) ₂
39	6	-CH ₂ Ph	-CH ₂ Ph-4-Cl	C(Me) ₂
40	5	-CH ₂ Ph-4-Cl	-S(O) ₂ -t-Bu	C(Me) ₂
41	5	-CH ₂ Ph-4-Cl	-S(O)-t-Bu	C(Me) ₂
42	6	-CH ₂ CHCH ₂	-CH ₂ Ph-4-Cl	C(Me) ₂

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TABLE II (cont.)

5	Ex No.	ATTACH POINT	R ⁸	R ⁵	Y-(CR ¹¹ R ¹¹) _p
	43	6	-(CH ₂) ₂ CH ₃	-CH ₂ Ph-4-Cl	C(Me) ₂
10	44	6	-CH ₂ CH ₃	-CH ₂ Ph-4-Cl	C(Me) ₂
	45	5	-CH ₂ Ph-4-Cl	-C(O)Ph-4-t-Bu	C(Me) ₂
	46	5	-CH ₂ Ph-4-Cl	-C(O)Ph-4-Cl	C(Me) ₂
	47	5	-CH ₂ Ph-4-Cl	-t-Bu	C(Me) ₂
15	48	5	-CH ₂ Ph-4-Cl	-C(O)Me	C(Me) ₂
	49	5	-CH ₂ Ph-4-Cl	-C(O)-c-Pr	C(Me) ₂
	50	5	-CH ₂ Ph-4-Cl	-C(O)CH ₂ CH ₂ -c-C ₅ H ₉	C(Me) ₂
20	51	5	-CH ₂ Ph-4-Cl	-C(O)CH ₂ CH(Me) ₂	C(Me) ₂
	52	5	-CH ₂ Ph-4-Cl	-C(O)Et	C(Me) ₂
	53	5	-CH ₂ Ph-4-Cl	-C(O)CH(Me) ₂	C(Me) ₂
	54	5	-CH ₂ Ph-4-Cl	-C(O)C(Me) ₃	C(Me) ₂
25	55	5	-CH ₂ Ph-4-Cl	-C(O)CH ₂ Ph	C(Me) ₂
	56	5	-CH ₂ Ph-4-F	-C(O)CH ₂ -t-Bu	C(Me) ₂
	57	5	-CH ₂ Ph-4-Br	-C(O)CH ₂ -t-Bu	C(Me) ₂
30	58	5	-CH ₂ Ph-4-I	-C(O)CH ₂ -t-Bu	C(Me) ₂
	59	5	-CH ₂ Ph-4-Cl	-C(Me) ₂ Pr	C(Me) ₂
	60	5	-CH ₂ Ph-4-Cl	-C(Me) ₂ Et	C(Me) ₂
	61	5	-CH ₂ Ph-3-F	-t-Bu	C(Me) ₂
35	62	5	-CH ₂ Ph-4-Cl	-CH(Me) ₂	C(Me) ₂
	63	5	-CH ₂ Ph-4-Cl	-c-Pr	C(Me) ₂
	64	5	-CH ₂ Ph-4-Cl	-(1-Me)-c-Pr	C(Me) ₂
	65	5	-CH ₂ Ph-4-Cl	-c-C ₅ H ₉	C(Me) ₂
40	66	5	-CH ₂ Ph-4-Cl	-c-C ₆ H ₁₁	C(Me) ₂
	67	5	-CH ₂ Ph-4-Cl	-C(Me) ₂ Ph	C(Me) ₂
	68	5	-CH ₂ Ph-4-Cl	-C(Me) ₂ Ph-4-Cl	C(Me) ₂
45	69	5	-CH ₂ Ph-4-Cl	-1-Ad	C(Me) ₂
	70	5	-CH ₂ Ph-4-Cl	-CH ₂ -1-Ad	C(Me) ₂
	71	6	-t-Bu	-CH ₂ Ph-4-Cl	C(Me) ₂
	72	6	-C(Me) ₂ Et	-CH ₂ Ph-4-Cl	C(Me) ₂
50	73	5	-CH ₂ Ph-4-Cl	-C(O)CH ₂ -t-Bu	C(Et) ₂

Assays for Determining Biological Activity

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Compounds of formula I can be tested using the following assays to determine their mammalian leukotriene biosynthesis inhibiting activity.

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Rat Peritoneal Polymorphonuclear (PMN) Leukocyte Assay

Rats under ether anesthesia are injected (i.p.) with 8 mL of a suspension of sodium caseinate (6 grams in ca. 50 mL water). After 15-24 hr. the rats are sacrificed (CO₂) and the cells from the peritoneal cavity are recovered by lavage with 20 mL of buffer (Eagles MEM containing 30 mM HEPES adjusted to pH 7.4 with NaOH). The cells are pelleted (350 x g, 5 min.), resuspended in buffer with vigorous shaking, filtered through lens paper, recentrifuged and finally suspended in buffer at a concentration of 10 cells/mL. A 500 mL aliquot of PMN suspension and test compound are preincubated for 2 minutes at 37 °C, followed by the addition of 10 mM A-23187. The suspension is stirred for an additional 4 minutes then bioassayed for LTB₄ content by adding an aliquot to a second 500 mL portion of the PMN at 37 °C. The LTB₄ produced in the first incubation causes aggregation of the second PMN, which is measured as a change in light transmission. The size of the assay aliquot is chosen to give a submaximal transmission change (usually -70%) for the untreated control. The percentage inhibition of LTB₄ formation is calculated from the ratio of transmission change in the sample to the transmission change in the compound-free control.

Human Polymorphonuclear (PMN) Leukocyte LTB₄ Assay

A. Preparation of Human PMN. Human blood was obtained by antecubital venepuncture from consenting volunteers who had not taken medication within the previous 7 days. The blood was immediately added to 10% (v/v) trisodium citrate (0.13 M) or 5% (v/v) sodium heparin (1000 IU/mL). PMNs were isolated from anticoagulated blood by dextran sedimentation of erythrocytes followed by centrifugation through Ficoll-Hypaque (specific gravity 1.077), as described by Boyum (Scand. J. Clin. Lab. Invest., 21 (Supp. 97), 77(1968)). Contaminating erythrocytes were removed by lysis following exposure to ammonium chloride (0.16 M) in Tris buffer (pH 7.65), and the PMNs resuspended at 5 x 10⁵ cells/mL in HEPES (15 mM)-buffered Hanks balanced salt solution containing Ca²⁺ (1.4 mM) and Mg²⁺ (0.7 mM), pH 7.4. Viability was assessed by Trypan blue exclusion and was typically greater than 98%.

B. Generation and Radioimmunoassay of LTB₄. PMNs (0.5 mL; 2.5 x 10⁵ cells) were placed in plastic tubes and incubated (37 °C, 2 min) with test compounds at the desired concentration or vehicle (DMSO, final concentration 0.2%) as control. The synthesis of LTB₄ was initiated by the addition of calcium ionophore A23187 (final concentration 10 mM) or vehicle in control samples and allowed to proceed for 5 minutes at 37 °C. The reactions were then terminated by the addition of cold methanol (0.25 mL) and samples of the entire PMN reaction mixture were removed for radioimmunoassay of LTB₄.

Samples (50 mL) of authentic LTB₄ of known concentration in radioimmunoassay buffer (RIA) buffer (potassium phosphate 1 mM; disodium EDTA 0.1 mM; Thimerosal 0.025 mM; gelatin 0.1%, pH 7.3) or PMN reaction mixture diluted 1:1 with RIA buffer were added to reaction tubes. Thereafter [³H]-LTB₄ (10 nCi in 100 mL RIA buffer) and LTB₄-antiserum (100 mL of a 1:3000 dilution in RIA buffer) were added and the tubes vortexed. Reactants were allowed to equilibrate by incubation overnight at 4 °C. To separate antibody-bound from free LTB₄, aliquots (50 mL) of activated charcoal (3% activated charcoal in RIA buffer containing 0.25% Dextran T-70) were added, the tubes vortexed, and allowed to stand at room temperature for 10 minutes prior to centrifugation (1500 x g; 10 min; 4 °C). The supernatants containing antibody-bound LTB₄ were decanted into vials and Aquasol 2 (4 mL) was added. Radioactivity was quantified by liquid scintillation spectrometry. Preliminary studies established that the amount of methanol carried into the radioimmunoassay did not influence the results. The specificity of the antiserum and the sensitivity of the procedure have been described by Rokach et al. (Prostaglandins Leukotrienes and Medicine 1984, 13, 21.) The amount of LTB₄ produced in test and control (approx. 20 ng/10⁶ cells) samples were calculated. Inhibitory dose-response curves were constructed using a four-parameter algorithm and from these the IC₅₀ values were determined.

Asthmatic Rat Assay

Rats are obtained from an inbred line of asthmatic rats. Both female (190-250 g) and male (260-400 g) rats are used.

Egg albumin (EA), grade V, crystallized and lyophilized, is obtained from Sigma Chemical Co., St. Louis. Aluminum hydroxide is obtained from the Regis Chemical Company, Chicago. Methysergide bimalate was supplied by Sandoz Ltd., Basel.

The challenge and subsequent respiratory recordings are carried out in a clear plastic box with internal dimensions (10 x 6 x 4 inches) (10 x 6 x 4) x 2.54 cm. The top of the box is removable; in use, it is held firmly in place by four clamps and an airtight seal is maintained by a soft rubber gasket. Through the center

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of each end of the chamber a Devilbiss nebulizer (No. 40) is inserted via an airtight seal and each end of the box also has an outlet. A Fleisch No. 0000 pneumotachograph is inserted into one end of the box and coupled to a Grass volumetric pressure transducer (PT5-A) which is then connected to a Beckman Type R Dynograph through appropriate couplers. While aerosolizing the antigen, the outlets are open and the pneumotachograph is isolated from the chamber. The outlets are closed and the pneumotachograph and the chamber are connected during the recording of the respiratory patterns. For challenge, 2 mL of a 3% solution of antigen in saline is placed into each nebulizer and the aerosol is generated with air from a small Potter diaphragm pump operating at (10 psi) 10 x 6,895 Pa and a flow of 8 liters/minute.

Rats are sensitized by injecting (subcutaneously) 1 mL of a suspension containing 1 mg EA and 200 mg aluminum hydroxide in saline. They are used between days 12 and 24 postsensitization. In order to eliminate the serotonin component of the response, rats are pretreated intravenously 5 minutes prior to aerosol challenge with 3.0 mg/kg of methysergide. Rats are then exposed to an aerosol of 3% EA in saline for exactly 1 minute, then their respiratory profiles are recorded for a further 30 minutes. The duration of continuous dyspnea is measured from the respiratory recordings.

Compounds are generally administered either orally 1-4 hours prior to challenge or intravenously 2 minutes prior to challenge. They are either dissolved in saline or 1% methocel or suspended in 1% methocel. The volume injected is 1 mL/kg (intravenously) or 10 mL/kg (orally). Prior to oral treatment rats are starved overnight. Their activity is determined in terms of their ability to decrease the duration of symptoms of dyspnea in comparison with a group of vehicle-treated controls. Usually, a compound is evaluated at a series of doses and an ED₅₀ is determined. This is defined as the dose (mg/kg) which would inhibit the duration of symptoms by 50%.

The invention is further defined by reference to the following examples, which are intended to be illustrative and not limiting. All temperatures are in degrees Celsius.

25 Example 1

3-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2, 2-dimethylpropanoic acid

Step A: 3-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-methoxyindol-2-yl]-2,2-dimethylpropanoic acid methyl ester

To a solution of 39 g of methyl 5-(t-butylthio)-2,2-dimethyl-4-oxopentanoate in a mixture of 300 mL of toluene and 150 mL of glacial acetic acid was added 15 g of NaOAc and 50 g of 1-(4-methoxyphenyl)-1-(p-chlorobenzyl)hydrazine hydrochloride. The reaction was maintained with stirring at room temperature for 3 days under argon in the dark. The mixture was poured into 3 L of H₂O and extracted with 3 x 500 mL of EtOAc. The ethyl acetate was washed with 3 x 500 mL of water then solid NaHCO₃ was added. The mixture was filtered and the filtrate washed twice with water. The organic phase was dried over MgSO₄ and evaporated to dryness to provide the title compound. m.p. 102-103 ° C.

Step B: 3-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-methoxyindol-2-yl]-2,2-dimethylpropanoic acid

The compound from Step A was hydrolysed using 325 mL of THF, 600 mL of MeOH and 325 mL of 1.0M LiOH. The solution was heated to 80 ° C for 3 h. The solution was acidified with 1N HCl and extracted with 3 x 200 mL of EtOAc. The organic phase was washed with water (2 x 150 mL) and dried over MgSO₄. The solution was evaporated to dryness to provide the title compound. m.p. 190-191 ° C.

Anal C, H, N:	Calc.	C 65.27;	H 6.57;	N 3.04,
	Found	C 65.28;	H 6.58;	N 3.04

Step C: Methyl 3-[N-(p-chlorobenzyl)-5-hydroxy-3-(t-butylthio)indol-2-yl]-2,2-dimethyl-propanoate

A solution of 61 mL of t-butylthiol in 650 mL of dry HMPA at 0 ° C was treated portionwise with 26 g of 50% NaH in mineral oil after removal of oil with hexane. The reaction was stirred at RT for 30 mins and 46 g of the compound from Step B was added.

The reaction was then heated under N₂ at 175 ° C for 5 hours. The solution was cooled, and poured onto crushed ice, after which it was treated with 2 N HCl to pH 5 and extracted with EtOAc (3 x 500 mL). The organic phase was washed with H₂O (3 x 200 mL) dried (MgSO₄) and evaporated. The residue was

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dissolved in 300 mL of ether and ethereal diazomethane was added until all acid was consumed. The excess solvent was removed and the oily residue triturated with hexane to leave a crystalline mass which was recrystallized from EtOAc/hexane to provide the title compound as a white crystalline solid, m.p. 170-171 ° C. From the mother liquors was isolated methyl 3-[N-(p-t-butylthiobenzyl)-5-hydroxy-3-(t-butylthio)-indol-2-yl]-2,2-dimethyl propanoate which was used as such in Example 3.

Step D: Methyl 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate.

Methyl 3-[N-(p-chlorobenzyl)-5-hydroxy-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoate (33.6 g) from Step C was dissolved in 500 mL of dry DMF and the solution was charged with 2.4 g of KI, 30.3 g of K₂CO₃, 4.77 g of Cs₂CO₃ and 23.5 g of 2-(chloromethyl)quinoline hydrochloride. The reaction was stirred at RT, under N₂, for 72 hours then it was poured into water (1.5 L), acidified with 1N HCl and extracted (3 x 200 mL) with CH₂Cl₂. The organic phase was washed with H₂O (3 x 150 mL), dried and evaporated. The residue was dissolved in hot EtOAc and upon cooling crystallized to deposit 22.0 g of the title compound, m.p. 166-167 ° C..

Step E: 3-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2, 2-dimethylpropanoic acid

Using the hydrolytic procedure of Step B but substituting the ester of Step D for the ester of Step A provided the title compound, which was recrystallized from 1:1 EtOAc/hexane. m.p. 208 ° C.

Anal C, H, N:	Calc. Found	C 69.55;	H 6.01;	N 4.77,
		C 69.77;	H 6.05;	N 4.70

Example 1A

3-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

Step A: N-Acetyl-4-(quinolin-2-ylmethoxy)aniline

A mixture containing 2-(chloromethyl)quinoline hydrochloride (100.0 g), 4-acetamidophenol (70.69 g) and milled anhydrous potassium carbonate (194 g) was stirred in DMF (1.2 L) using a mechanical stirrer for 48 hours. The mixture was carefully poured onto ice/water (3 L) with vigorous stirring. After the ice had melted, the solid was filtered and rinsed thoroughly with water. It was recrystallized from 95% ethanol and filtered to give the title compound in three crops.

Step B: 4-(Quinolin-2-ylmethoxy)aniline

A suspension of N-acetyl-4-(quinolin-2-ylmethoxy)aniline (Step A, 108.9 g) in 1 L of 95% ethanol containing 10 M KOH (120 mL) was heated at reflux under nitrogen in a heating mantle. When the hydrolysis was complete (approx. 36 h), the reaction mixture was cooled and ethanol was partially removed under vacuum. The mixture was then diluted with water (200 mL) and the fine off-white crystals were collected and thoroughly rinsed with water. The material, after air-drying, yielded the title compound which was used as such in the next step.

Step C: 4-(Quinolin-2-ylmethoxy)phenylhydrazine

A quantity of 84 g of 4-(quinolin-2-ylmethoxy)aniline from Step B was suspended in 300 mL of deionized H₂O and 84 mL of 12 M HCl. The suspension was stirred vigorously to obtain a fine particle suspension. Then a precooled solution (5 ° C) of 23.88 g of sodium nitrite dissolved in 75 mL of deionized H₂O was added dropwise to the suspension at 5 ° C over 25 minutes. The solution was stirred at 5 ° C for 60 min to obtain the diazonium salt as a clear brown solution. The presence of excess HNO₂ was confirmed by KI-starch paper, and the pH of the solution was about 3.0. If a white suspension persisted after 1 h, the mixture was filtered through a glass wool plug, to give the diazonium salt in the filtrate.

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In the meantime a sodium hydrosulfite solution was prepared by dissolving 321 g of sodium hydrosulfite (approx. 85% purity) in 2 L of deionized water, and cooled at 0° to 5°C. To this solution were added 15 mL of 2N NaOH and 2 L of ether. The biphasic solution was kept near 0°C by addition of crushed ice and was stirred vigorously. To this solution was added dropwise the diazonium salt solution with stirring maintained throughout. At the end of the addition an orange solid was formed and 600 mL of NaOH (2N) was added over 30 minutes. The reaction was finally stirred for 60 minutes at 25°C. The solid was collected, suspended in ether (1 L) and filtered. The process was repeated with 2 L of water to yield the title compound as a pale yellow solid after freeze-drying overnight. m.p. 73-85°C (dec).

Step D: 1-(p-Chlorobenzyl)-1-[4-(quinolin-2-yl-methoxy)phenyl]hydrazine

A quantity of 10 g of 4-(quinolin-2-ylmethoxy)phenylhydrazine from Step C was added to a solution of 10.5 mL of diisopropylethylamine and 150 mL of CH₂Cl₂. To the yellow suspension was added 9.11 g of p-chlorobenzyl chloride followed by 3.64 g of Bu₄NBr and 50 mL of CH₂Cl₂. The reaction was stirred for approximately 24 hours. When no starting material remained, the reaction was diluted with H₂O and extracted 3 times with CH₂Cl₂. The combined organic phase was washed once with water and dried (MgSO₄), filtered and evaporated to dryness. The solid residue was dried under vacuum overnight prior to being swished in ether/methanol 90/10 to give the title compound as a pale yellow solid. m.p. 130°C.

Step E: 3-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The methyl ester of the title compound was prepared according to the method described in Step A of Example 1 but using the phenylhydrazine from Step D of Example 1A as starting material.

The title compound was prepared under the conditions described in Step B of Example 1.

Example 2

3-[N-(p-Chlorobenzyl)-3-methyl-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared according to the method of Example 1, but using methyl 2,2-dimethyl-4-oxohexanoate as starting material in Step A in place of methyl 5-t-butylthio-2,2-dimethyl-4-oxopentanoate. m.p. 215-217°C.

Example 3

3-[N-(p-t-Butylthiobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The methyl ester byproduct from Step C of Example 1 was reacted 2-(chloromethyl)quinoline according to the conditions of Steps D & E of Example 1 to provide the title compound. m.p. 172-173°C.

Example 4

3-[N-(p-Chlorobenzyl)-3-(phenylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared according to the method described for Example 1, but substituting methyl-5-phenylthio-2,2-dimethyl-4-oxopentanoate for methyl 5-t-butylthio-2,2-dimethyl-4-oxopentanoate in Example 1 (Step A).

Anal. C, H, N for sodium salt• 2 H ₂ O:			
Calc.	C 64.91;	H 5.30;	N 4.20
Found	C 64.94;	H 5.04;	N 4.15

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Example 5

3-[N-(p-Chlorobenzyl)-3-(phenylsulfonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid, N-oxide

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Methyl 3-[N-(p-chlorobenzyl)-3-(phenylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethyl propanoate (430 mg) from Step D of Example 4 was dissolved in 5 mL cold CH₂Cl₂ and treated with a solution of 448 mg of 80% m-chloroperbenzoic acid (MCPBA) in CH₂Cl₂. After 24 hours, the solution was poured onto 10 mL of sat. aqueous NaHCO₃ solution, extracted with 3 x 10 mL of CH₂Cl₂, washed with 2 x 10 mL of H₂O, dried with magnesium sulfate and evaporated to dryness. The residue was crystallised from 2:1 CH₂Cl₂/EtOAc to yield 280 mg of the title compound as its methyl ester. Hydrolysis using the conditions described in Example 1 (Step B) provided the title compound, m.p. 197 ° C (dec.)

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Anal. C, H, N:	Calc.	C 66.0;	H 4.77;	N 4.28
	Found	C 66.06;	H 4.77;	N 4.19.

Examples 6 and 7

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3-[N-(p-Chlorobenzyl)-3-(phenylsulfonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid and 3-[N-(p-Chlorobenzyl)-3-(phenylsulfinyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

Methyl 3-[N-p-chlorobenzyl-3-(phenyl thio)-5-(quinolin-2-methoxy)indol-2-yl]-2,2-dimethyl propanoate (430 mg) from Example 4 (Step D) was dissolved in 5 mL of cold methylene chloride and a solution of 150 mg of 80% (MCPBA) in methylene chloride was added. After 24 hours, the reaction solution was poured onto 10 mL of saturated aqueous sodium bicarbonate solution and this mixture was extracted 3 times with 10 mL of methylene chloride. The combined organic phases were washed twice with 10 mL of water, dried with magnesium sulfate and evaporated under vacuum.

Chromatography over silica gel (2 hexane: 1 ethyl acetate) provided two compounds which were separately hydrolyzed using the procedure described in Example 1 (Step B).

3-[N-(p-Chlorobenzyl)-3-(phenylsulfonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid:

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Anal. C, H, N for sodium salt • H ₂ O:			
Calc.	C 63.57;	H 4.89;	N 4.12
Found	C 63.28;	H 4.77;	N 3.90

3-[N-(p-Chlorobenzyl)-3-(phenylsulfinyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid:

40

Anal. C, H, N for sodium salt • H ₂ O:			
Calc.	C 63.38;	H 5.17;	N 4.11
Found	C 63.28;	H 4.89;	N 3.97

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Example 8

3-[N-(p-Chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

Step A: Methyl 3-[N-(p-chlorobenzyl)-5-hydroxyindol-2-yl]-2,2-dimethylpropanoate

A suspension of 1.0 g of 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-methoxyindol-2-yl]-2,2-dimethylpropanoic acid (from Example 1 Step B) in 50 mL of CH₂Cl₂ was treated with 1.3 mL of ethanethiol and 3.47 g of AlCl₃ at 0 ° C under argon. After 40 min the mixture was poured onto 50 mL 1N HCl, extracted with 3 x 50 mL of CH₂Cl₂ washed with 2 x 50 mL of H₂O, dried with MgSO₄ and the solvent removed. The residue was dissolved in 10 mL ether and ethereal diazomethane added until all the acid was consumed.

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The excess solvent was removed and the residue chromatographed on silica gel to afford the title compound.

Step B: 3-[N-(p-Chlorobenzyl)-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared by treating the ester from Step A with 2-(chloromethyl)quinoline hydrochloride under the conditions of Step D and effecting hydrolysis under the conditions of Example 1 (Step B), m.p. 193-194 °C.

Example 9

3-[N-(p-Chlorobenzyl)-3-benzoyl-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid

Step A: Methyl 3-[N-(p-chlorobenzyl)-3-benzoyl-5-benzoyloxyindol-2-yl]-2,2-dimethylpropanoate

Methyl 3-[N-(p-chlorobenzyl)-5-hydroxy indol-2-yl]-2,2-dimethylpropanoate (609 mg) from Example 8 (Step A) was dissolved in 10 mL of 1,2-dichloroethane and the solution charged with 0.5 mL of benzoyl chloride and 680 mg of AlCl₃. The reaction was heated to 80 °C under argon for 1.5 h, then quenched with 20 mL of 0.5N Na, K tartrate solution, extracted with 3 x 20 mL of ether, washed with 10 mL of H₂O and dried (MgSO₄). Removal of solvent provided an oily residue which was chromatographed on silica gel to give the title compound.

Step B: Methyl 3-[N-(p-chlorobenzyl)-3-benzoyl-5-hydroxyindol-2-yl]-2,2-dimethylpropanoate

The compound from Step A (300 mg) was dissolved in 4 mL of MeOH and treated with 1 mL of a 1.4 M solution of NaOMe in MeOH under argon for 3 hrs. The mixture was poured onto 20 mL of NH₄ OAc (25% solution), extracted with 3 x 15 mL of ether, washed with 10 mL of H₂O, dried over MgSO₄ and the solvent removed under vacuum. The resulting oil was purified by chromatography on silica gel to afford the title compound.

Step C: 3-[N-(p-Chlorobenzyl)-3-benzoyl-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared using the conditions described in Step D and Step E of Example 1, but substituting the ester from Step B for the ester of Example 1, Step C; m.p. 165-166 °C.

Example 10

3-[N-(p-Chlorobenzyl)-3-benzyl-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid

Step A: Methyl 3-[N-(p-chlorobenzyl)-3-benzyl-5-(benzoyloxy)indol-2-yl]-2,2-dimethylpropanoate

Methyl 3-[N-(p-chlorobenzyl)-3-benzoyl-5-(benzoyloxy)indol-2-yl]-2,2-dimethylpropanoate (360 mg) (prepared in Step A of Example 9), 800 mg of ZnI₂, and 500 mg of sodium cyanoborohydride were stirred in 5 mL of dichloroethane at RT under argon for 30 min. The temperature was then raised to 65 °C for 3 hr. After the solution had cooled, it was poured onto 10 mL of NH₄ OAc (25% solution), extracted with 3 x 15 mL of ether, washed with 10 mL of H₂O and dried (MgSO₄). The solution was evaporated to dryness and the residue was chromatographed on silica gel to yield the title compound as a white foam.

Step B: 3-[N-(p-Chlorobenzyl)-3-benzyl-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared under the conditions described in Step B and Step C of Example 9 but substituting the ester from Example 10 (Step A) for the ester of Example 9 (Step A), m.p. 178 °C.

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Example 11

3-[N-(p-Chlorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2, 2-dimethyl-
propanoic acid

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The title compound was prepared according to the method described in Example 9, but using t-butylacetylchloride in place of benzoyl chloride in Step A, m.p. 183-184 °C.

Example 12

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2-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]ethoxyethanoic acid

Step A: Methyl 2-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl] ethanoate

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The title compound was prepared according to the method outlined in Steps A-D of Example 1, but using methyl 4-t-butylthio-3-oxo-butanoate in Step A instead of methyl 5-t-butylthio-2,2-dimethyl-4-oxopentanoate.

Step B: 2-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]ethanol

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The compound from Step A (192 mg) was dissolved in 3 mL of THF at RT under an argon atmosphere and treated with 30 mg of lithium aluminum hydride. After 1 hr, the reaction was poured onto 10 mL of 0.5 N Na,K tartrate solution and extracted with 3 x 10 mL of EtOAc. The organic layer was washed with 10 mL of H₂O, dried (MgSO₄) and evaporated to dryness to yield the title compound.

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Step C: 2-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]ethoxyethanoic acid

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To 91 mg of 2-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]ethanol from Step B in 2 mL THF at 0 °C under an argon atmosphere was added 40 mg of 80% sodium hydride over 30 min. Ethyl bromoacetate (0.3 ml) was added to the solution and the reaction stirred at RT overnight. The reaction was poured onto 10 mL of NH₄OAc (25% solution), extracted with 3 x 10 mL of EtOAc, washed with 20 mL of H₂O and dried over MgSO₄. Removal of the solvent followed by column chromatography on silica gel afforded the ethyl ester of title compound. Hydrolysis of this ester under the conditions described in Step B of Example 1 provided the title compound, m.p. 185 °C (dec.).

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Example 13

3-[N-(p-Chlorobenzyl)-3-(3,3-dimethyl-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

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The title compound was prepared according to the method described in Example 10 but using methyl 3-[N-(p-chlorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(t-butylacetyloxy)indol-2-yl]-2,2-dimethylpropanoate (obtained as an intermediate from Example 11) as starting material, m.p. 188 °C (dec.).

Example 14

45

3-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2-methylpropanoic acid

50

The title compound was prepared according to the method of Example 1 using methyl 5-t-butylthio-2-methyl-4-oxopentanoate as starting material in Step A in place of methyl 5-t-butylthio-2,2-dimethyl-4-oxopentanoate.

¹H NMR (250 MHz, acetone-d₆) δ 1.05 (3H, d, J = 6Hz), 1.15 (9H, s), 2.7 (1H, m), 3.2 (2H, d, J = 7Hz), 5.4 (2H, s), 5.6 (2H, s), 6.9 (1H, dd), 7.0 (2H, d), 7.3 (4H, m), 7.6 (1H, td), 7.7 (1H, d), 7.8 (1H, td), 7.9 (1H, d), 8.1 (1H, d), 8.3 ppm (1H, d).

55

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Example 15

3-[N-(p-Chlorobenzyl)-3-methyl-5-(6,7-dichloroquinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

5 The title compound was prepared according to the method described in Example 1 but using methyl 2,2-dimethyl-4-oxohexanoate as starting material in Step A and 2-(bromomethyl)-6,7-dichloroquinoline in Step D.

10

Anal. C, H, N:	Calc.	C 63.21;	H 4.74;	N 4.91
	Found	C 63.47;	H 4.94;	N 4.67

Example 16

15

3-[N-(p-Chlorobenzyl)-3-methyl-5-(7-chloroquinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared according to the method described in Example 15 but using 2-(bromomethyl)-7-chloroquinoline instead of 2-(bromomethyl)-6,7-dichloroquinoline. m.p. 105-107 °C.

20

Anal. C, H, N:	Calc.	C 67.41;	H 5.24;	N 5.24
	Found	C 67.82;	H 5.12;	N 4.32

25

Example 17

3-[N-(p-Chlorobenzyl)-4-allyl-5-(quinolin-2-ylmethoxy)-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoic acid30 Step A: Methyl 3-[N-(p-chlorobenzyl)-5-allyloxy-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoic acid

500 mg. of methyl 3-[N-(p-chlorobenzyl)-5-hydroxy-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoate from Step C of Example 1 was dissolved in 5 mL of DMF and 20 mg of K₂CO₃ and 150 mg of allyl bromide were added. The reaction was stirred for 16 hrs. Water was added and the organic phase extracted with EtOAc (3 x 5mL). The organic phase was dried with MgSO₄ and evaporated to yield, after chromatography on silica gel (EtOAc:hexane 1:5), the title compound.

35

Step B: Methyl 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-4-allyl-5-hydroxyindol-2-yl]-2,2-dimethylpropanoate

40 500 mg of the ester of Step A was converted to the title compound by heating to 180 ° in m-xylene for 4 hours.

Step C: 3-[N-(p-Chlorobenzyl)-4-allyl-5-(quinolin-2-ylmethoxy)-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoic acid

45

The title compound was prepared from the compound of Step B using the methodology of Example 1, (Steps D and E), m.p. 103-105 °C.

50

Anal. C, H, N:	Calc.	C 69.09;	H 6.11;	N 4.35
	Found	C 70.55;	H 6.31;	N 4.29

55

EP 0 419 049 B1Example 183-[N-(p-Chlorobenzyl)-4-allyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

5 The methyl ester of the title compound was prepared according to the method of Example 17 but substituting methyl 3-[N-(p-chlorobenzyl)-5-hydroxyindol-2-yl]-2,2-dimethylpropanoate as starting material (obtained in Step A Example 8) for the ester in Example 17 (Step A). Hydrolysis was then effected according to the conditions of Step B of Example 1 to provide the title compound, m.p. 196-197 °C (dec.).

10 Example 193-[N-(p-Chlorobenzyl)-6-(quinolin-2-ylmethoxy)-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoic acid

15 The title compound was prepared according to the conditions of Example 1, Steps A to E, but substituting 1-(3-methoxyphenyl)-1-(p-chlorobenzyl)hydrazine hydrochloride for the starting material in Example 1 (Step A). Chromatographic separation of the desired regioisomer was achieved at Step A by isolating the most polar product, methyl 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-6-methoxyindol-2-yl]-2,2-dimethylpropanoate. The properties of the title compound were as follows: m.p. 165-167 °C.

20

Anal C, H, N:	Calc.	C 69.54;	H 6.01;	N 4.77
	Found	C 69.46;	H 6.18;	N 4.96

25 Example 203-[N-(p-Chlorobenzyl)-4-(quinolin-2-ylmethoxy)-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoic acid

30 Methyl 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-4-methoxyindol-2-yl]-2,2-dimethylpropanoate was obtained as a by-product from Step A of Example 19 and isolated by chromatography as the less polar product. The compound was used as starting material for the preparation of the title product using the methodology of Steps B to E of Example 1.

35

Anal C, H, N:	Calc.	C 69.54;	H 6.01;	N 4.77;
	Found	C 69.80;	H 6.24;	N 4.86

Example 21

40

3-[N-(p-Chlorobenzyl)-3-(t-butylthio)-7-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

45 The title product was prepared according to Steps A to E of Example 1 but substituting 1-(2-methoxyphenyl)-1-(p-chlorobenzyl)hydrazine hydrochloride for 1-(4-methoxyphenyl)-1-(p-chlorobenzyl)hydrazine hydrochloride in Example 1 (Step A), m.p. 206 °C.

50

Anal. C, H, N:	Calc.	C 69.54;	H 6.01;	N 4.77,
	Found	C 69.40;	H 5.88;	N 4.65

55

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Example 22

2-[2-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]ethoxy]propanoic acid sodium salt dihydrate

5

Step A: Methyl 2-[2-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]ethoxy]propanoate

The title compound was prepared from 251 mg of 2-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]ethanol (Step B of Example 12) under the conditions described in Step C of Example 12 using methyl D,L-2-bromopropanoate instead of ethyl bromoacetate.

10

Step B: 2-[2-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]ethoxy]propanoic acid sodium salt dihydrate

The acid corresponding to the title compound of Example 22 was prepared from the ester of Step A of Example 22 under the conditions described in Step B of Example 1. A quantity of 204 mg of the acid was suspended in 1.5 mL of EtOH and treated with 1 equiv. of 1N aq. NaOH and freeze dried for 2 days to afford the title compound.

20

Anal. C, H, N:	Calc.	C 61.25;	H 5.61;	N 4.33,
	Found	C 61.75;	H 5.70;	N 3.97

25 Example 23.

3-[N-(p-Chlorobenzyl)-4-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

Step A: Methyl 3-[N-(p-chlorobenzyl)-4-hydroxyindol-2-yl]-2,2-dimethylpropanoate

30

The title compound was prepared using methodology from Step A of Example 8 but substituting 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-4-methoxy indol-2-yl]-2,2-dimethylpropanoic acid (Step B of Example 20) for the propanoic acid in Example 8 (Step A).

Step B: 3-[N-(p-Chlorobenzyl)-4-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title product was prepared according to conditions described in Steps D and E of Example 1 substituting methyl 3-[N-(p-chlorobenzyl)-4-hydroxy indol-2-yl]-2,2-dimethylpropanoate for the propanoate in Example 1 (Step D), m.p. 158-160 °C.

40

Anal. C, H, N:	Calc.	C 72.20;	H 5.45;	N 5.61
	Found	C 72.25;	H 5.60;	N 5.75

45

Example 24

3-[N-Methyl-3-(p-chlorobenzoyl)-6-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid

Step A: Methyl 3-[6-methoxy-3-(t-butylthio) indol-2-yl]-2,2-dimethylpropanoate

A mixture of 4.2 g of 3-methoxyphenylhydrazine hydrochloride and 4.9 g of methyl 5-(t-butylthio)-2,2-dimethyl-4-oxopentanoate in 100 mL of t-butanol was refluxed for 18 hours. The mixture was cooled to R.T., and evaporated to dryness. The residue was suspended in ether (150 ml) and stirred for 30 min. The salts were filtered and the filtrate evaporated to dryness to give a residue which was chromatographed on flash silica gel using as eluant ethyl acetate:toluene (1:99) to isolate the title compound as the most polar product; m.p. 133 °C.

55

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Step B: Methyl 3-[N-methyl-3-(t-butylthio)-6-methoxyindol-2-yl]-2,2-dimethylpropanoate

A solution of 1.75 g of the indole from Step A in 30 mL of THF and 3 mL HMPA was cooled to -78 °C and to this solution was slowly added a solution of 0.54M KHMDS in toluene (10.2 mL). The mixture was stirred at this temperature for 15 min. and treated with 0.34 mL of iodomethane. The mixture was stirred at -78 °C for 5 h, quenched with 1N HCl (100 mL), extracted with ethyl acetate, and the organic layer washed with H₂O, dried over Na₂SO₄ and evaporated to dryness. The residue was chromatographed on flash silica gel using ethyl acetate:hexane (20:80) as eluant to afford the title compound as a solid; m.p. 97-98 °C.

Step C: Methyl 3-[N-methyl-6-hydroxyindol-2-yl]-2,2-dimethylpropanoate

To a cold solution of 940 mg of the indole ester from Step B and 1.6 mL of ethanethiol in CH₂Cl₂ (50 mL) was added portion-wise 4.3 g of AlCl₃. After complete addition, the mixture was stirred at R.T. for 2 h. The mixture was then cooled to 0 °C and carefully quenched with a solution of 0.5 M Na,K tartrate (200 mL) and extracted with CH₂Cl₂. The organic layer was dried over Na₂SO₄ and evaporated to dryness to give a solid which was chromatographed on flash silica gel using ethyl acetate:hexane (30:70) as eluant to afford the title compound; m.p. 125-126 °C.

Step D: Methyl 3-[N-methyl-6-(p-chlorobenzoyloxy)-3-(p-chlorobenzoyl)indol-2-yl]-2,2-dimethylpropanoate

To a cold solution of 393 mg of hydroxy indole from Step C in 5 mL of THF were added 0.31 mL of Et₃N followed by 0.21 mL of p-chlorobenzoyl chloride. The mixture was stirred at R.T. for 15 min and quenched with H₂O. The mixture was extracted with ethyl acetate which was dried over Na₂SO₄ and evaporated to dryness to give a solid which was dissolved in 10 mL of 1,2-dichloroethane. To this mixture were added successively at R.T. 0.38 mL of p-chlorobenzoyl chloride and 803 mg of AlCl₃. The mixture was heated at 80 °C for 3 h, cooled to R.T. and quenched with 50 mL of 0.5 N HCl. The mixture was extracted with CH₂Cl₂, washed with H₂O, dried over Na₂SO₄ and evaporated to dryness. The residue was chromatographed on flash silica gel using ethyl acetate:hexane (20:80) as eluant to afford the title compound as a white solid. m.p. 138 °C.

Step E: Methyl 3-[N-methyl-3-(p-chlorobenzoyl)-6-hydroxyindol-2-yl]-2,2-dimethylpropanoate

To a suspension of 270 mg of the p-chlorobenzoate from Step D in 3 mL of MeOH was added 1.2 mL of a solution of 1.3M NaOMe in MeOH and the mixture was stirred at R.T. for 2 h. The reaction mixture was poured onto 25% aq. NH₄OAc and extracted with ethyl acetate. The organic extract was dried over Na₂SO₄, evaporated to dryness and the residue chromatographed on flash silica gel using ethyl acetate:hexane (40:60) as eluant to afford the title compound as a yellow foam.

Step F: Methyl 3-[N-methyl-3-(p-chlorobenzoyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate

To a solution of 180 mg of the phenol from Step E in 5 mL of DMF were added 124 mg of milled K₂CO₃ followed by 150 mg of 2-(bromomethyl) quinoline. The mixture was stirred at R.T. for 18 h, poured onto 25% aq. NH₄OAc and extracted with ethyl acetate. The extract was dried over Na₂SO₄ and evaporated to dryness to give an oil which was chromatographed on flash silica gel using ethyl acetate:hexane (30:70) as eluant to give the title compound as a foam.

Step G: 3-[N-methyl-3-(p-chlorobenzoyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

To a solution of 230 mg of ester from Step F in 1.5 mL of THF and 3 mL of MeOH was added 1M aq. LiOH and the mixture stirred at 80 °C for 4 h. The mixture was cooled to R.T. and evaporated to dryness in vacuo. The residue was dissolved in a mixture of 20 mL of 25% aq. NH₄OAc and 20 mL of ethyl acetate using vigorous stirring. The organic layer was separated, dried over Na₂SO₄ and evaporated to dryness to give a yellow solid (216 mg). This solid was swished for 2 h in 5 mL of a mixture of Et₂O:hexane (1:1). The solid was filtered and rinsed with a 1:2 mixture of Et₂O:hexane to give the title product as a yellow solid, m.p. 203-205 °C.

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Anal. C, H, N:	Calc.	C 70.65;	H 5.16;	N 5.32;
	Found	C 70.42;	H 5.25;	N 5.40

5

Example 25

3-[N-methyl-3-(p-chlorobenzyl)-6-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid, sodium salt hemihydrate

10

Step A: Methyl 3-[N-methyl-3-(p-chlorobenzyl)-6-(p-chlorobenzoyloxy)indol-2-yl]-2,2-dimethylpropanoate

To a solution of 500 mg of the benzoyl derivative from Step D of Example 24 in 10 mL of 1,2-dichloroethane were added 1.19 g of ZnI₂ and 700 mg of NaBH₃CN. The mixture was heated at 65 °C for 5
15 hours and cooled to R.T. The mixture was quenched with 1N aq. HCl and extracted with CH₂Cl₂. The extracts were washed with brine, dried over Na₂SO₄ and evaporated to dryness to give an oil which was chromatographed on flash silica gel using ethyl acetate:hexane (15:85) as eluant to isolate the title compound as a white foam.

Step B: Methyl 3-[N-methyl-3-(p-chlorobenzyl)-6-hydroxyindol-2-yl]-2,2-dimethylpropanoate

To a suspension of 425 mg of p-chlorobenzoate from Step A in 3 mL of MeOH was added 1.9 mL of a solution of 1.3M NaOMe in MeOH. The mixture was stirred at R.T. for 1 h, poured into 20 mL of 25% aq. NH₄OAc, and extracted with ethyl acetate. The organic extract was dried over Na₂SO₄ and evaporated to
25 dryness to give an oil which was chromatographed on flash silica gel using ethyl acetate:hexane (30:70) as eluant to give the title compound as a white foam.

Step C: Methyl 3-[N-methyl-3-(p-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate

To a solution of 315 mg of the ester from Step B in 3 mL of DMF were added 225 mg of milled K₂CO₃ and 272 mg of 2-(bromomethyl)quinoline. The mixture was stirred at R.T. for 18 h, poured into 25% aq. NH₄OAc, and extracted with ethyl acetate. The organic extract was dried over Na₂SO₄ and evaporated to
30 dryness to give an oil which was chromatographed on flash silica gel using ethyl acetate:hexane (30:70) as eluant to give the title compound as a foam.

35

Step D: 3-[N-methyl-3-(p-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethyl propanoic acid, sodium salt hemihydrate

To a solution of 367 mg of the ester from Step C in 3 mL of THF and 6 mL of MeOH was added 1 M
40 aq. LiOH and the mixture was heated at 80 °C for 2 h. The mixture was cooled to R.T. and evaporated to dryness. The residue was dissolved in a mixture of 20 mL of 25% aq. NH₄OAc and 20 mL of ethyl acetate (vigorous stirring required). The organic layer was separated, dried over Na₂SO₄ and evaporated to dryness to give a white solid (346 mg). The solid was swished at R.T. for 2 h with 10 mL of a mixture of Et₂O:hexane (1:1), filtered, rinsed with a mixture of (1:2) Et₂O:hexane and the solid collected to give the title
45 compound as its free acid, a white solid: m.p. 185 °C.

The title compound was prepared by dissolving the above acid in 1 mL of EtOH to which 0.63 mL of 1N aq. NaOH was added. The mixture was freeze dried for 2 days to give the title product as a white solid.

50

Anal. C, H, N:	Calc.	C 67.32;	H 5.29;	N 5.07;
	Found	C 67.15;	H 5.35;	N 5.17

55

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EXAMPLE 263-[N-(4-Chlorobenzyl)-3-i-propoxy-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

5 The title compound is prepared according to the method of Example 1, but using methyl 5-i-propoxy-2,2-dimethyl-4-oxopentanoate as starting material in Step A in place of methyl 5-t-butylthio-2,2-dimethyl-4-oxopentanoate.

EXAMPLE 27

10

3-[N-(4-Chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2-ethylpropanoic acidStep A: Methyl 4-chloro-2-ethyl-4-pentenoate

15 A 2L 3-necked flask equipped with a mechanical stirrer, pressure equalizing addition funnel, and nitrogen inlet was charged with diisopropylamine (28 mL, 200 mmol) and dry THF (400 mL). The mixture was cooled to 0 °C and a 1.6 M solution of butyl lithium in hexane (125 mL, 200 mmol) was then added over a 15 minute period and stirring was continued for an additional 45 minutes.

The resultant solution of lithium diisopropylamide (200 mmol) was cooled to -78 °C and then butyric acid (9.1 mL, 100 mmol) was added over a 15 minute period. The reaction was allowed to warm to room temperature (1 hour) and then heated at 55 °C for 3.5 hours. The resultant gel was cooled to -78 °C and then treated 2,3-dichloro-1-propene (10.1 mL, 110 mmol) over a 15 minute period. The mixture was then allowed to warm to room temperature and stirred for 18 hours.

25 The reaction mixture was diluted with Et₂O (400 mL), extracted with H₂O (400 mL) and with NaOH 1N (300 mL). The aqueous layers were combined, acidified with HCl (2N, until pH 1-2) and the product was extracted with EtOAc (2 x 300 mL). The organic layers were combined, washed with brine (200 mL) and dried over MgSO₄. Filtration and concentration gave a yellow oil which was dissolved in dry MeOH (150 mL) and acetyl chloride (1 mL) was added dropwise. The resultant solution was gently refluxed for 20 hours. The reaction was allowed to cool to room temperature and it was concentrated. The resultant residue was diluted with Et₂O (600 mL), washed with NaHCO₃ sat. (200 mL), washed with brine (200 mL), and dried over MgSO₄. Filtration and concentration gave a yellow oil which was purified by Kugelrohr distillation (bp 110 °C at 0.2 mm Hg) to give pure (250 MHz NMR) methyl 4-chloro-2-ethyl-4-pentenoate.

Step B: Methyl 5-bromo-2-ethyl-4-oxopentanoate

35

To a cold (0 °C) solution of methyl 4-chloro-2-ethyl-4-pentenoate from Step 2 (1.67 g, 9.5 mmol) in MeOH (31 mL) and H₂O (16 mL) was added Br₂ dropwise (0.60 mL, 11.6 mmol). The resulting yellow solution was stirred at room temperature for 1 hour. EtOAc (300 mL) and H₂O (200 mL) were added. The organic layer was separated, washed with H₂O, 1N NaOH, H₂O, brine and dried over MgSO₄. Filtration and concentration gave a yellow liquid which was purified by flash chromatography (EtOAc/Hexane (1:9)) to give pure (NMR 250 MHz) methyl 5-bromo-2-ethyl-4-oxopentanoate.

Step C: Methyl 5-(t-butylthio)-2-ethyl-4-oxopentanoate

45

To a cold (0 °C) stirred solution of the bromoketone from Step 2 (490 mg, 2.07 mmol), in 10 mL of dry THF, were sequentially added 2-methyl-2-propyl thiol (0.30 mL) and triethylamine (0.40 mL, 2.9 mmol). The reaction mixture was then allowed to warm to room temperature. After 18 hours the white solid was removed by filtration and the filtrate was concentrated. The resulting yellow residue was purified by flash chromatography (Et₂O/Hexane (9:50)) to give the pure title compound (250 MHz NMR).

50

Step D: 3-[N-(4-Chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-yl-methoxy)indol-2-yl]-2-ethylpropanoic acid.

The title compound was prepared according to the method of Example 1, but using methyl 5-t-butylthio-2-ethyl-4-oxopentanoate as starting material in Step A in place of methyl 5-t-butylthio-2,2-dimethyl-4-oxopentanoate.

55

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Anal. C, H, N, for sodium salt • 2H ₂ O:			
Calc.	C 63.30;	H 5.94;	N 4.34
Found.	C 63.29;	H 5.87;	N 4.37

5

The sodium salt of the title compound in this and other Examples was prepared by the method of Example 25.

10 EXAMPLE 28

3-[N-(4-Chlorobenzyl)-3-trifluoroacetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

Step A: Methyl 3-[N-(4-chlorobenzyl)-3-trifluoroacetyl-5-hydroxyindol-2-yl]-2,2-dimethylpropanoate

15 Methyl 3-[N-(4-chlorobenzyl)-5-hydroxy indol-2-yl]-2,2-dimethylpropanoate (310 mg) from Example 8 (Step A) was dissolved in 3 ml of 1,2-dichloroethane and the solution charged with 0.6 ml of trifluoroacetic anhydride and 500 mg of AlCl₃. The reaction was stirred at RT, under argon for 4h, then quenched with 20 mL of 0.5 N Na, K tartrate solution, extracted with 3x20 ml of Et₂O, washed with 10 mL of H₂O and dried over MgSO₄. Removal of solvent provided an oily residue which was chromatographed on silica gel to give
20 the title compound.

Step B: 3-[N-(4-Chlorobenzyl)-3-trifluoroacetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid.

25 The title compound was prepared using the conditions described in Step D and Step E of Example 1, but substituting the ester from Step A for the ester of Example 1, Step C.

Anal. C, H, N, for sodium salt • 7H ₂ O			
Calc.	C 51.72;	H 5.29;	N 3.76
Found.	C 51.81;	H 5.19;	N 3.73

30

35 EXAMPLE 29

3-[N-(4-Chlorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2-methylpropanoic acid

40 Step A: Methyl 5-(t-butylthio)-2-methyl-4-oxopentanoate

The title compound was prepared according to the method described in Example 27, but using propionic acid as starting material in Step A in place of butyric acid.

45 Step B: 3-[N-(4-chlorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2-methylpropanoic acid

The title compound was prepared according to the method described in Example 11, but substituting 5-t-butylthio-2-methyl-4-oxopentanoate as starting material in Example 1 Step A in place of methyl 5-t-butylthio-2,2-dimethyl-4-oxopentanoate.
50

Anal. C, H, N, for sodium salt • 1.5 H ₂ O			
Calc.	C 66.50;	H 5.90;	N 4.43
Found.	C 66.58;	H 5.87;	N 4.40

55

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EXAMPLE 30

3-[3-(3,3-Dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid5 Step A: Methyl 3-[3-(t-butylthio)-5-methoxy indol-2-yl]-2,2-dimethylpropanoate

A mixture containing 4-methoxyphenyl hydrazine HCl (70.66 g, 0.405 mol) and methyl 5-(t-butylthio)-2,2-dimethyl-4-oxopentanoate (99.54 g, 0.405 mol) in ^tBuOH (400 ml) was heated at a gentle reflux for 48 hours. The mixture was allowed to cool to RT and the precipitated NH₄Cl was removed by filtration. The residue was concentrated and fractionated on a plug of silica using EtOAc/hexane (1:3) as eluent. Evaporation of the appropriate fraction gave an orange-brown solid which was crystallized from EtOH (100 ml). Yield from two crops afforded 55.6g of the title compound.
¹H NMR (CD₃COCD₃): δ 1.20(s, 6H); 1.25 (s, 9H) 3.33 (s, 2H); 3.62 (s, 3H); 3.81 ppm (s, 3H); in addition to aromatic protons.

15 Step B: Methyl 3-[5-hydroxyindol-2-yl]-2,2-dimethylpropanoate

To a solution of the compound from step A (25.50 g, 73 mmol) in CH₂Cl₂ (250 ml) at 0 °C was added AlCl₃ (87.70 g, 9 mol eq.) portion-wise. When the addition was complete, the ice-bath was removed and the mixture was stirred at RT for 3 hours. EtSH (27 ml, 5 mol eq.) was added and the resulting mixture was stirred for a further 5 hours. It was then slowly poured onto an ice-cold 1M solution of Na, K tartrate. The product was extracted into CH₂Cl₂ (x2) and the organic phase was washed with aq. NaCl (x3). Conventional work-up followed chromatography on silica gel using EtOAc/hexane 1:5 to 3:2 afforded 13.60 g of the title compound, 75% yield.
¹H NMR (CDCl₃): δ 1.26 (s, 6H); 2.95 (s, 2H); 3.72 (s, 3H); 6.11 (s, 1H); 6.71 (bd, 1H); 6.95 (s, 1H); 7.16 (d, 1H); 7.26 ppm (s, 1H)

Step C: Methyl 3-[5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate

A mixture of the phenol from step B (13.62 g, 55.14 mmol) and 2-bromomethylquinoline (12.85 g, 1.05 mol eq.) and anhydrous K₂CO₃ (15.22 g, 2 mol. eq.) in DMF (40 ml) was stirred at RT for 48 hours. The mixture was then poured onto ice/water and after all the ice had melted, the brown solid was collected and air-dried. The dried material was passed through a plug of silica (using EtOAc/hexane (1:3) as eluent) to remove the color; yield: 19 g, 88%.
Recrystallization from EtOH afforded 14.17g of pure title compound, m.p. 131-132 °C.

Step D: Methyl 3-[3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate

To a suspension of AlCl₃ (5.7 g, 42 mmol) in CH₂Cl₂ (30 mL) at 0 °C was added 3,3-dimethylbutanoyl chloride (2.4 mL, 17 mmol). After 15 minutes at 0 °C, a solution of the ester from step C (3.0 g, 7.7 mmol) in CH₂Cl₂ (10 mL) was added by double-tipped needle. The mixture was stirred a further 20 minutes, at which point it was poured into a mixture of 0.5 M Na, K tartrate (150 mL) and ice (100 g). The product was extracted with EtOAc, and the organic layer was washed successively with 0.5 M NaK tartrate, H₂O, and brine. The solvent was then removed and yellow/orange oil (3.5 g) was used without purification in the following step.

Step E: 3-[3-(3,3-Dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The crude ester from step D was dissolved in a mixture of MeOH (20 mL), THF (20 mL), and H₂O (5 mL). To this was added 10 M NaOH (2.3 mL, 23 mmol). After stirring for 3 hours, the solution was cooled to 0 °C, and HOAc (1.5 mL) was added dropwise. The solution was partly concentrated to remove the THF and MeOH, and the product was then extracted into EtOAc. The organic layer was washed with H₂O and brine. After drying (MgSO₄), the solution was filtered and evaporated to give a pale orange solid. The product was stirred vigorously with a mixture of isopropanol (30 mL) and H₂O (3 mL) to give the title compound as an off-white amorphous solid (2.6 g). mp = 193-196 °C (dec)

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EXAMPLE 31

3-[N-(4-Trifluoromethylbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid

5 The product from Example 30 (100 mg, 0.21 mmol) and 4-trifluoromethylbenzyl bromide (98 mg, 0.41 mmol), and methyltriocetylammmonium chloride (83 mg, 0.21 mmol) were dissolved in a mixture of 50% NaOH (2 mL) and benzene (0.5 mL). After vigorous stirring for 3.5 hours, the reaction mixture was cooled to 0 °C and was acidified with HOAc (2 mL). The product was extracted with EtOAc, and the organic layer
10 was washed with H₂O and brine. Following evaporation of the solvent, the residue was purified by flash chromatography on silica gel, eluting with 1:5 EtOAc/hexane containing 1% HOAc. The resulting yellow foam was triturated with 1:4 Et₂O/hexane to give the title compound as a pale yellow solid (37 mg, 28%).
¹H NMR (CDCl₃) δ 8.22 (1H, d, J=8.5 Hz), 8.13 (1H, d, J=8.5 Hz), 7.87-7.70 (3H, m), 7.62-7.45 (4H, m), 7.00-6.85 (4H, m), 5.45 (4H, s), 3.58 (2H, brs), 2.87 (2H, s), 1.30 (6H, s), 1.03 ppm (9H, s).

EXAMPLE 32

3-[N-Benzyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

20 Following the method of Example 31, with benzyl bromide as the alkylating agent, the title compound was obtained as an off-white solid (mp = 180-183 °C (dec)).

EXAMPLE 33

3-[N-(3-Methoxybenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

25 Following the method of example 31, with 3-methoxybenzyl bromide as the alkylating agent, the title compound was obtained as an off-white solid. (mp = 173-175 °C (dec)).

EXAMPLE 34

3-[N-Allyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

35 Step A: Allyl 3-[N-allyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate

To a solution of the product from Example 30 (100 mg, 0.21 mmol) in dry DMF (2 mL) was added 80% NaH (14 mg, 0.47 mmol), followed 15 minutes later by allyl bromide (0.35 mL, 4 mmol). After 2 1/2 hours,
40 saturated NH₄Cl solution was added, and the product was extracted with EtOAc. The organic layer was washed with H₂O and brine, dried over MgSO₄, filtered and was then evaporated to give the title compound as a yellow oil which was used as such in the next step.

45 Step B: 3-[N-Allyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The crude ester from step 1 was treated as in the method of Example 30, Step 2 to give the title compound as an off-white solid (mp = 146-148 °C (dec)).

EXAMPLE 35

3-[N-(4-Methoxybenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

55 Following the method of Example 34, with 4 methoxybenzyl chloride as the alkylating agent, the title compound was obtained as a white solid.
¹H NMR (250 MHz, acetone-d₆) δ 1.05 (9H, s), 1.27 (6H, s), 2.38 (2H, s), 3.73 (3H, s), 3.78 (2H, s), 5.46 (2H, s), 5.48 (2H, s), 6.70-8.40 ppm (13H, aromatics).

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EXAMPLE 36

3-[N-Methyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

5 Following the method of Example 34, with methyl iodide as the alkylating agent, the title compound was obtained as a white solid.

Anal. C, H, N for sodium salt . 1 H ₂ O			
Calc.	C 68.42;	H 6.70;	N 5.32
Found.	C 68.36;	H 6.81;	N 5.44

EXAMPLE 37

3-[3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acidStep A: 3-(Quinolin-2-ylmethoxy)phenylhydrazine

20 The title compound was prepared using the conditions described in Step A, Step B and Step C of Example 1A, but replacing the phenol in Step A with 3-acetamidophenol; m.p. 55-70 °C (dec).

Step B: Ethyl (4-chlorobenzene)propanoate

25 A solution of 4-chlorobenzaldehyde (28 g) and (carboethoxymethylene)triphenylphosphorane (73 g) in toluene (500 mL) was refluxed for 2 hours. The reaction was cooled to room temperature and concentrated under vacuum to a total volume of 150 mL. Then pure hexane (500 mL) was added and the mixture was left for 18 hours at room temperature. The solid (triphenylphosphine oxide) was filtered, rinsed with hexane and the filtrate evaporated to give crude product which was distilled at 0.5 mm Hg, and the fraction boiling at
30 130 °C was collected to give ethyl 4-chlorocinnamate, which was reduced as follows: the cinnamate (21 g) was hydrogenated in EtOAc (300 mL) in the presence of 5% Pd on C (2 g) for 3 hours at atmospheric pressure. After completion, the reaction mixture was filtered through a celite pad, rinsed with EtOAc and the filtrate evaporated to dryness to give the title product as an oil.

35 Step C: Methyl 6-(4-chlorophenyl)-2,2-dimethyl-4-oxohexanoate

To a solution of ethyl (4-chlorobenzene)propanoate (Step B, 10 g) in dry THF (500 mL) at -78 °C was added 0.58M potassium hexamethyldisilazane in toluene (243 mL). The mixture was stirred at -60 °C for one hour. Then a solution of 2,2-dimethylsuccinic anhydride (6 g) in THF (100 mL) was added slowly and
40 the mixture was slowly warmed to room temperature and finally stirred for 18 hours. Water (1000 mL) was added and the organic layer separated. The aqueous layer was washed with EtOAc (3 x 250 mL) and acidified with 1N HCl. The aqueous layer was extracted with EtOAc, the extract was dried (Na₂SO₄) and evaporated to give a residue which was dissolved in THF (100 mL) and MeOH (200 mL) and treated at reflux with 1N LiOH (100 mL) for 4 hours. The mixture was cooled to room temperature and concentrated
45 under vacuum until H₂O distilled off. Water (500 mL) was added, the mixture acidified with 1N aq. HCl, extracted with EtOAc, the extract was dried (Na₂SO₄) and evaporated to give the title product as its carboxylic acid. The compound was treated with diazomethane in ether, evaporated to dryness and chromatographed over silica gel, eluting with EtOAc-hexane (10:90) to give the title product as a white solid;
50 m.p. 52-54 °C.

Step D: Methyl 3-[3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate

55 To a solution of methyl 6-(4-chlorophenyl)-2,2-dimethyl-4-oxohexanoate (Step C, 2.8 g) in toluene (30 mL) and glacial HOAc (15 mL) was added portion-wise solid 3-(quinolin-2-ylmethoxy)phenylhydrazine (Step A, 3.2 g) and stirred at room temperature for 2 hours. The reaction mixture was diluted with Et₂O (200 mL), washed with 1N NaOH, H₂O, dried (Na₂SO₄) and evaporated to give crude hydrazone which was immediately treated as follows: the crude hydrazone was dissolved in a mixture of PPE (15 mL) and 1,2-dichloroethane (30 mL) and stirred at 40 °C for 18 hours. The reaction mixture was cooled to 0 °C and

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carefully treated with 1N NaOH to bring to pH 9. The mixture was then extracted with ether, the extract was washed with H₂O, dried (Na₂SO₄) and evaporated to give a residue, which was chromatographed in a column of flash silica gel (eluting with EtOAc-hexane 25:75) and isolating the most polar component as the title product as a foam.

Step E: 3-[3-(4-Chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared using the conditions described in Step B of Example 1, but substituting the ester from Step D for the ester of Example 1, Step A.

Anal. C, H, N for sodium salt • 1 1/2 H ₂ O			
Calc.	C 65.75;	H 5.33;	N 5.11
Found.	C 66.08;	H 5.31;	N 5.08

EXAMPLE 38

3-[N-(Phenylsulfonyl)-3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

Step A: Methyl 3-[N-(phenylsulfonyl)-3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2 -yl]-2,2-dimethylpropanoate

To a solution of methyl 3-[3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate (Step D, Example 37) (208 mg) in dry THF (5 mL) and HMPA (0.5 mL) at -78°C was added 0.58M potassium hexamethyldisilazane in toluene (0.77 mL) and the solution then stirred at -78°C for 15 minutes. Then freshly distilled benzenesulfonyl chloride (0.062 mL) was added and stirred at -78°C for 2.5 hours. The reaction mixture was quenched with 25% aq. NH₄OAc, extracted with EtOAc, dried over Na₂SO₄ and evaporated to give crude product. Chromatography of the residue in a column of flash silica gel (eluting with EtOAc-hexane 25:75) afforded the title product as an oil.

Step B: 3-[N-Phenylsulfonyl)-(3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethyl propanoic acid

The title compound was prepared under the conditions described in Step B of Example 1, but substituting the ester from Step A for the ester of Example 1, Step A.

Anal. C, H, N for sodium salt			
Calc.	C 67.65;	H 4.89;	N 4.38
Found.	C 68.07;	H 5.18;	N 4.32

EXAMPLE 39

3-[N-Benzyl-3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared under the conditions described in Step A and Step B of Example 38, but substituting benzyl chloride for the benzenesulfonyl chloride from Example 38 (Step A).

Anal. C, H, N for sodium salt • 1/2 H ₂ O			
Calc.	C 71.66;	H 5.36;	N 4.52
Found.	C 71.65;	H 5.49;	N 4.44

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EXAMPLE 40 AND 41

3-[N-(4-Chlorobenzyl)-3-(t-butylsulfonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid and
 3-[N-(4-chlorobenzyl)-3-(t-butylsulfonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

5

The title compounds were prepared using the conditions described for Examples 6 and 7, but substituting the ester of Example 4 (Step D) for the ester of Example 1 (Step D).

3-[N-(4-chlorobenzyl)-3-(t-butylsulfonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid:

10

Anal. C, H, N for sodium salt • 1 1/2 H ₂ O			
Calc.	C 61.12;	H 5.58;	N 4.19
Found	C 61.31;	H 5.39;	N 4.19

15

3-[N-(4-chlorobenzyl)-3-(t-butylsulfonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid:

20

Anal. C, H, N for sodium salt • 2 H ₂ O			
Calc.	C 61.76;	H 5.79;	N 4.24
Found	C 61.44;	H 5.68;	N 4.19

EXAMPLE 42

25

3-[N-Allyl-3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared under the conditions described in Step A and Step B of Example 38, but substituting allyl bromide for the benzenesulfonyl chloride from Example 38 (Step A).

30

Anal. C, H, N for sodium salt • 2 H ₂ O			
Calc.	C 66.38;	H 5.74;	N 4.69
Found	C 66.57;	H 5.75;	N 4.73

35

EXAMPLE 43

40

3-[N-(n-Propyl)-3-(4-chlorobenzyl)-6-(quinoline-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

A solution of methyl 3-[N-allyl-3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate (Example 42, methyl ester) (190 mg) was hydrogenated in EtOAc (4 mL) in the presence of 5% Pd on charcoal at atmospheric pressure for 1 hour. Filtration on Celite pad and evaporation of liquors afforded the methyl ester of the title product. Hydrolysis of this ester under the conditions described in Step B of Example 1 provided the title compound.

45

Anal. C, H, N for sodium salt • 1 1/2 H ₂ O			
Calc.	C 67.17;	H 5.98;	N 4.75
Found	C 67.38;	H 5.44;	N 4.85

50

55

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EXAMPLE 44

3-[N-Ethyl-3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

5 The title product was prepared under the conditions described in Step A and Step B of Example 38, but substituting iodoethane for the benzenesulfonyl chloride from Example 38 (Step A).

Anal. C, H, N for sodium salt . 2 H ₂ O			
Calc.	C 65.69;	H 5.86	N 4.79
Found	C 65.81;	H 5.21;	N 4.77

EXAMPLE 45

3-[N-(4-Chlorobenzyl)-3-(4-t-butylbenzoyl)-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid

15 The title compound was prepared according to the method described in Example 9, but using 4-t-butylbenzoyl chloride in place of benzoyl chloride in Step A.

Anal. C, H, N, for sodium salt . 2 1/2 H ₂ O			
Calc.	C 67.81;	H 5.97;	N 3.86
Found.	C 67.91;	H 6.01;	N 3.64

EXAMPLE 46

3-[N-(4-chlorobenzyl)-3-(4-chlorobenzoyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

30 The title compound was prepared according to the method described in Example 9, but using 4-chlorobenzoyl chloride in place of benzoyl chloride in Step A.

Anal. C, H, N for sodium salt . 2 H ₂ O			
Calc.	C 63.89;	H 4.78;	N 4.03
Found	C 64.20;	H 4.61;	N 3.99

EXAMPLE 47

3-[N-(4-Chlorobenzyl)-3-(1,1-dimethylethyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid45 Step A: Methyl 3-[N-(4-chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate

Methyl 3-[N-(4-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate (3.77 g, 6.27 mmol) from Step D of Example 1, was dissolved in 75 mL of dry CH₂Cl₂ and the solution was charged with 6.27 g (47.0 mmol) of AlCl₃ and the mixture was stirred at RT, under Ar, for 1.75 hours. The reaction was then quenched by the addition of 0.5N Na, K tartrate (150 mL) and the resulting mixture was extracted with EtOAc (3x). The organic extracts were washed with 0.5N Na, K tartrate (1x) and with brine (1x), and dried (MgSO₄). Filtration and removal of solvents provided a brown oily residue which was chromatographed on silica gel using EtOAc-hexane (1:3) to give the title compound.

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Step B: Methyl 3-[N-(4-chlorobenzyl)-3-(1,1-dimethylethyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate

Trimethylacetyl chloride (4.23 g, 4.32 mL, 35.09 mmol) was added to a cold suspension (0 °C) of AlCl₃ (11.7 g, 87.7 mmol) in dry CH₂Cl₂ (60 mL) under Ar. The yellow mixture was stirred at 0 °C for 15 minutes, and a solution of 9.00 g (17.54 mmol) of methyl 3-[N-(4-chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate (prepared in Step A) in CH₂Cl₂ (40 mL) was added dropwise (over 10 minutes) at 0 °C. The reaction mixture was stirred for 10 minutes and slowly poured onto an ice-cold and vigorously stirred mixture of 0.5M aqueous Na, K tartrate (500 mL) and EtOAc (400 mL). After 20 minutes, the aqueous layer was extracted with EtOAc (2x) and the combined organic extracts were washed with H₂O (2x), with 1N aqueous NaOH (2x), with H₂O (2x) and dried (MgSO₄). Filtration and removal of solvents provided a yellow oily residue which was chromatographed on silica gel using EtOAc-hexane (1:4) to give the title compound.

Step C: 3-[N-(4-chlorobenzyl)-3-(1,1-dimethylethyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared according to the conditions described in Step B of Example 1, but substituting the ester from Step B for the ester of Example 1. The title compound was recrystallized from EtOAc-EtOH; m.p. 201-202 °C.

Anal. Calc.	C 73.57	H 6.36	N 5.05
Found	C 73.75	H 6.37	N 5.03

EXAMPLE 483-[N-(4-Chlorobenzyl)-3-acetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared according to the conditions described in Step B and Step C of Example 47, from methyl 3-[N-(4-chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate (prepared in Step A of Example 47) but using acetyl chloride in place of trimethylacetyl chloride in Step B. ¹H NMR (CD₃COCD₃) δ 1.20 (6H, s), 2.64 (3H, s), 3.62 (2H, br s), 5.47 (2H, s), 5.57 (2H, s), 6.90-6.99 (3H, m), 7.28-7.37 (3H, m), 7.56-7.83 (4H, m), 7.97 (1H, d), 8.05 (1H, d), 8.39 ppm (1H, d).

EXAMPLE 493-[N-(4-Chlorobenzyl)-3-cyclopropanecarbonyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared according to the conditions described in Step B and Step C of Example 47, from methyl 3-[N-(4-chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate (prepared in Step A of Example 47), but using cyclopropanecarbonyl chloride in place of trimethylacetyl chloride in Step B.

Anal. C, H, N for sodium salt • 1 H ₂ O:			
Calc.	C 67.27;	H 5.31;	N 4.61
Found	C 67.27;	H 5.16;	N 4.58

EXAMPLE 503-[N-(4-chlorobenzyl)-3-(3-cyclopentylpropanoyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared according to the conditions described in Step B and Step C of Example 47, from methyl 3-[N-(4-chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate

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(prepared in Step A of Example 47), but using 3-cyclopentylpropanoyl chloride in place of trimethylacetyl chloride in Step B.

¹H NMR (CD₃COCD₃): δ 1.09 (2H, m), 1.22 (6H, s), 1.40-1.91 (9H, m), 2.94 (2H, t), 3.68 (2H, br s), 5.46 (2H, s), 5.58 (2H, s), 6.91-6.99 (3H, m), 7.30 (3H, m), 7.53-7.63 (2H, m), 7.72-7.82 (2H, m), 7.96 (1H, d), 8.06 (1H, d), 8.34 ppm (1H, d).

EXAMPLE 51

10 3-[N-(4-Chlorobenzyl)-3-(3-methylbutanoyl)-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared according to the conditions described in Step B and Step C of Example 47, from methyl 3-[N-(4-chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethyl propanoate (prepared in Step A of Example 47), but using 3-methylbutanoyl chloride in place of trimethylacetyl chloride in Step B.

15 ¹H NMR (CD₃COCD₃): δ 0.98 (6H, d), 1.24 (6H, s), 2.30 (1H, m), 2.85 (2H, d), 3.70 (2H, br s), 5.46 (2H, s), 5.58 (2H, s), 6.96 (3H, m), 7.30 (3H, m), 7.55-7.63 (2H, m), 7.73-7.82 (2H, m), 7.95 (1H, d), 8.07 (1H, d), 8.36 ppm (1H, d).

EXAMPLE 52

20 3-[N-(4-Chlorobenzyl)-3-propanoyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared according to the conditions described in Step B and Step C of Example 47, from methyl 3-[N-(4-chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate (prepared in Step A of Example 47), but using propanoyl chloride in place of trimethylacetyl chloride in Step B.

30

Anal. C, H, N for sodium salt . 1H ₂ O			
Calc.	C 66.61;	H 5.42;	N 4.71
Found	C 66.87;	H 5.45;	N 4.69

35 EXAMPLE 53

3-[N-(4-chlorobenzyl)-3-(2-methylpropanoyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

40 The title compound was prepared according to the conditions described in Step B and Step C of Example 47, from methyl 3-[N-(4-chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate (prepared in Step A of Example 47), but using 2-methylpropanoyl chloride in place of trimethylacetyl chloride in Step B.

45 ¹H-NMR (CD₃COCD₃): δ 1.07 (6H, d), 1.16 (6H, s), 3.34 (1H, m), 3.64 (2H, br s), 5.46 (2H, s), 5.57 (2H, s), 6.95 (3H, m), 7.32 (3H, m), 7.45 (1H, br s), 7.60 (1H, br t), 7.71-7.83 (2H, m), 7.97 (1H, d), 8.07 (1H, d), 8.36 ppm (1H, d).

EXAMPLE 54

50 3-[N-(4-Chlorobenzyl)-3-trimethylacetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid, sodium salt

Step A: Methyl 3-[N-(4-chlorobenzyl)-3-trimethylacetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate

55 **Method A:** The indole from Example 47, Step A (2.00 g, 3.9 mmol) and trimethylacetyl chloride (0.86 g, 7.1 mmol) were dissolved in sieve-dried CH₂Cl₂ (15 mL). The mixture was cooled to -25 °C and AlCl₃ (1.64 g, 12.3 mmol) was added in two portions 5 minutes apart. After 10 minutes at -20 to -25 °C, 7 mL of 2.5 M aqueous HOAc was added to the mixture such that the temperature stayed below -20 °C. The mixture was then warmed to RT and the layers were separated. The organic layer was washed with H₂O, saturated

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aqueous NaHCO₃, and H₂O, and then evaporated to dryness. The resulting oil was crystallized from MeOH (20 mL) to give 1.4 g (60%) of the title compound.

Method B: A solution of TiCl₄ (6 mL of a 1.0 M solution in CH₂Cl₂, 6.0 mmol) and trimethylacetyl chloride (0.491 g, 4.1 mmol) was cooled to -5 °C. To the cooled solution was added a solution of the indole from Example 47, Step A (1.025 g, 2.0 mmol) in 2 mL of CH₂Cl₂ over a 5 minute period. After 30 minutes, the reaction was quenched by the addition of 3 mL of 2.5 M aqueous HOAc. The mixture was warmed to RT and the layers separated. The organic layer was washed with H₂O, saturated aqueous NaHCO₃, and H₂O, and then evaporated to dryness. The residual oil was crystallized from 10 mL of MeOH to give 625 mg (53%) of the title compound.

¹H-NMR (CDCl₃) δ 1.22 (6H, s), 1.30 (9H, s), 3.29 (2H, s), 3.61 (3H, s), 5.28 (2H, s), 5.47 (2H, s), 6.7-8.2 ppm (13H, m)

IR(Nujol mull) 1636, 1730 cm⁻¹

Step B: 3-[N-(4-Chlorobenzyl)-3-trimethylacetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid, sodium salt

The acylated methyl ester from Step A (401 mg, 0.67 mmol), absolute EtOH (1.62 g), and NaOH (64.2 mg of a 50.9% aq. solution, 0.82 mmol) were refluxed 42 hours. During the latter stages of the reaction, product crystallized. At the end of the reflux, the product was filtered and washed with EtOH to give 288 mg (74%) orange solid. The material was slurried in 3 mL EtOH for 8 hours at RT to give 190 mg of the title compound as a pale orange solid.

¹H-NMR (CD₃OD) δ 1.08 (6H, s), 1.15 (9H, s), 3.23 (2H, s), 5.40 (2H, s), 5.55 (2H, s), 6.7-8.2 ppm (13H, m). IR (Nujol mull) 1680, 1575 cm⁻¹.

EXAMPLE 55

3-[N-(4-Chlorobenzyl)-3-phenylacetyl-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid

The title compound was prepared according to the conditions described in Step B and Step C of Example 47, from methyl 3-[N-(4-chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate (prepared in Step A of Example 47), but using phenylacetyl chloride in place of trimethylacetyl chloride in Step B.

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Anal. C, H, N for sodium salt . 1H ₂ O			
Calc.	C 69.46;	H 5.22;	N 4.26
Found	C 69.71;	H 5.25;	N 4.11

EXAMPLE 56

3-[N-(4-Fluorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

Using the procedure of Example 31, but replacing 4-trifluoromethylbenzyl bromide with 4-fluorobenzyl bromide, the title compound is obtained.

EXAMPLE 57

3-[N-(4-Bromobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

Using the procedure of Example 31, but replacing 4-trifluoromethylbenzyl bromide with 4-bromobenzyl bromide, the title compound is obtained.

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EXAMPLE 58

3-[N-(4-Iodobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid

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Using the procedure of Example 31, but replacing 4-trifluoromethylbenzyl bromide with 4-iodobenzyl bromide, the title compound is obtained.

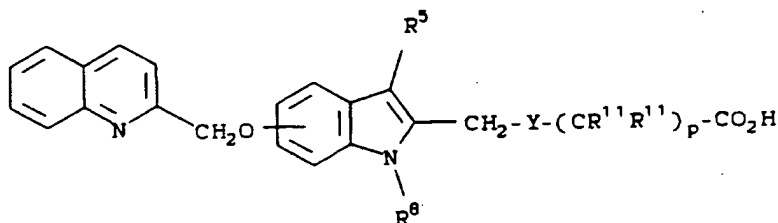
EXAMPLES 59-70

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Operating as described in the previous examples, the following compounds are prepared:

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Ex No.	ATTACH POINT	R ⁸	R ⁵	Y-(CR ¹¹ R ¹¹) _p
59	5	-CH ₂ Ph-4-Cl	-C(Me) ₂ Pr	C(Me) ₂
60	5	-CH ₂ Ph-4-Cl	-C(Me) ₂ Et	C(Me) ₂
61	5	-CH ₂ Ph-3-F	-C(Me) ₃	C(Me) ₂
62	5	-CH ₂ Ph-4-Cl	-CH(Me) ₂	C(Me) ₂
63	5	-CH ₂ Ph-4-Cl	-c-Pr	C(Me) ₂
64	5	-CH ₂ Ph-4-Cl	-(1-Me)-c-Pr	C(Me) ₂
65	5	-CH ₂ Ph-4-Cl	-c-C ₅ H ₉	C(Me) ₂
66	5	-CH ₂ Ph-4-Cl	-c-C ₆ H ₁₁	C(Me) ₂
67	5	-CH ₂ Ph-4-Cl	-C(Me) ₂ Ph	C(Me) ₂
68	5	-CH ₂ Ph-4-Cl	-C(Me) ₂ Ph-4-Cl	C(Me) ₂
69	5	-CH ₂ Ph-4-Cl	-1-Ad	C(Me) ₂
70	5	-CH ₂ Ph-4-Cl	-CH ₂ -1-Ad	C(Me) ₂
71	6	-t-Bu	-CH ₂ Ph-4-Cl	C(Me) ₂
72	6	-C(Me) ₂ Et	-CH ₂ Ph-4-Cl	C(Me) ₂

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EXAMPLE 73

3-[N-(4-Chlorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-diethylpropanoic acid

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Step A: Methyl 4-chloro-2,2-diethyl-4-pentenoate

To a cold solution (0 °C) of diisopropylamine (0.80 mL, 5.7 mmol) in THF (5 mL) was added a 1.6 M solution of butyllithium in hexane (3.4 mL, 5.4 mmol) over a 5 minute period and stirring was continued for an additional 45 min. Then a solution of methyl 4-chloro-2-ethyl-4-pentenoate (800 mg, 5 mmol) from Example 27, Step A, in THF (2 mL) was added and the reaction was stirred at 0 °C for another 30 minutes. Then ethyl iodide (440 µL, 5.5 mmol) was added and the reaction was allowed to proceed at room temperature for 2 hours. The reaction was quenched with NH₄OAc buffer (50 mL of 25% w/v) and extracted with EtOAc. The organic layer was separated, dried over MgSO₄, filtered and concentrated. The crude product was purified by Kugelrohr distillation (b.p. 120 °C at 0.1 mm Hg) to give the title compound.

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Step B: Methyl 5-(t-Butylthio)-2,2-diethyl-4-oxopentanoate

The title compound was prepared according to the method of Example 27, Step B and Step C, but using methyl 4-chloro-2,2-diethyl-4-pentenoate as starting material in Step B in place of methyl 4-chloro-2-ethyl-4-pentenoate.

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Step C: Methyl 3-[N-(4-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy) indol-2-yl]-2,2-diethylpropanoate.

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To a mixture of 1-(4-chlorobenzyl)-1-[4-(quinolin-2-ylmethoxy)phenyl]hydrazine from Example 1A, Step D, (660 mg, 1.7 mmol) and anhydrous NaOAc (160 mg, 1.95 mmol) in toluene (3 mL) was added glacial HOAc (1.5 mL). After 30 minutes, a solution containing methyl 5-(t-butylthio)-2,2-diethyl-4-oxopentanoate from Step B (402 mg, 1.47 mmol) in toluene (1 mL) was added and the reaction mixture stirred for 24 hours at room temperature and for 48 hours at 65 °C. The reaction was then diluted with EtOAc, washed with NH₄OAc buffer (25% w/v) and dried over MgSO₄. Filtration and concentration gave a viscous oil which was purified by flash chromatography on silica gel (eluant: EtOAc-hexane 15:85) to give the title compound.

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¹H NMR (250 MHz acetone-d₆): δ 0.85 (6H, t), 1.1 (9H, s), 1.7 (4H, q), 3.2 (2H, s), 3.6 (3H, s), 5.4 (2H, s), 5.5 (2H, s), 6.9 (3H, m), 7.3 (4H, m), 7.6 (1H, dd), 7.7 (1H, d), 7.8 (1H, td), 7.9 (1H, d), 8.1 (1H, d), 8.3 ppm (1H, d).

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Step D: Methyl 3-[N-(4-chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-diethylpropanoate

The title compound was prepared according to the method of Example 47, Step A, but using methyl 3-[N-(4-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-diethylpropanoate from Step C in place of methyl 3-[N-(4-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate.

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Step E: Methyl 3-[N-(4-chlorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-diethylpropanoate

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To a cold solution (0 °C) of methyl 3-[N-(4-chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-diethylpropanoate (from Step D) (177 mg, 0.33 mmol) in CH₂Cl₂ (3 mL) was added AlCl₃ (220 mg, 1.65 mmol) followed by t-butylacetyl chloride (82 µL, 0.66 mmol). The reaction was stirred at 0 °C for 20 minutes and then quenched with 30 mL of 0.5 N Na,K tartrate solution, and extracted with 3 x 30 mL of EtOAc. The organic layers were combined and dried over MgSO₄. Filtration and concentration gave an oily residue which was purified by flash chromatography (eluent EtOAc-hexane (17:83)) to give the title compound.

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Step F: 3-[N-(4-Chlorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy) -indol-2-yl]-2,2-diethylpropanoic acid

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The compound from Step E was hydrolysed using THF (2.5 mL), MeOH (0.6 mL) and NaOH (1N, 1.5 mL). The solution was heated at 70 °C for 2 weeks. The reaction was neutralized by addition of Na₄OAc

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buffer (20 mL of 25% w/v) and extracted with EtOAc (3 x 30 mL). The organic layers were combined, dried over MgSO₄, filtered and concentrated. The resulting residue was purified by flash chromatography (eluent: EtOAc-hexane-HOAc (250:750:1)) to give the title compound.

Anal. C, H, N for sodium salt • 1 1/2 H ₂ O;			
Calc.	C 67.70;	H 6.43;	N 4.15
Found	C 67.67;	H 6.31;	N 4.06

EXAMPLE 74

Methyl 3-[N-(4-chlorobenzyl)-3,6-bis(acetyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2 dimethylpropanoate

The title compound was isolated in Step B of Example 48 from a chromatography on silica gel (EtOAc-Hexane 2:3).

¹H NMR (CD₃COCD₃): δ 1.22 (6H, s), 2.63 (3H, s), 2.66 (3H, s), 3.58 (3H, s), 3.65 (2H, s), 5.61 (2H, s), 5.65 (2H, s), 6.95 (2H, d), 7.31 (2H, d), 7.62 (1H, br t), 7.71-7.85 (4H, m), 7.97 (1H, d), 8.08 (1H, d) 8.40 ppm (1H, d).

EXAMPLE 75

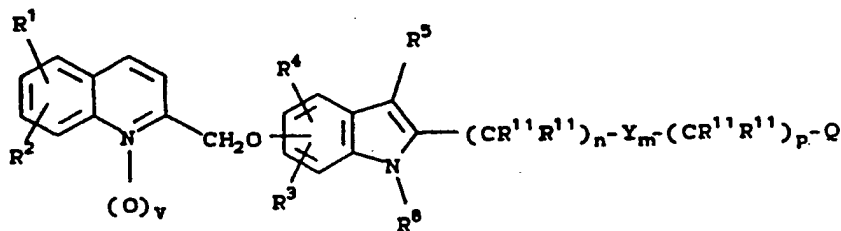
Methyl 3-[N-(4-chlorobenzyl)-3,6-bis (cyclopropanecarbonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoate

The title compound was isolated in Step B of Example 49 from a chromatography on silica gel (EtOAc-hexane 3:7) and was recrystallized from EtOAc-EtOH; m.p. 166-167 ° C.

Claims

Claims for the following Contracting States : AT, BE, CH, DE, DK, FR, GB, IT, LI, LU, NL, SE

1. A compound of the formula I:



I

wherein:

R¹, R², R³, R⁴ and R¹⁰ are independently hydrogen, halogen, C₁₋₇ alkyl, C₂₋₇ alkenyl, C₂₋₇ alkynyl, -CF₃, -CN, -NO₂, -N₃, -C(OH)R¹¹R¹¹, -CO₂R¹², -SR¹⁴, -S(O)R¹⁴, -S(O)₂R¹⁴, -S(O)₂NR¹⁵R¹⁵, -OR¹⁵, -NR¹⁵R¹⁵, -C(O)R¹⁵ or -(CH₂)₁R²¹;

R⁵ is hydrogen, -CH₃, -CF₃, -C(O)H, X¹-R⁶ or X²-R⁷;

R⁶ and R⁹ are independently: alkyl, -(CH₂)_uPh(R¹⁰)₂ or -(CH₂)_uTh(R¹⁰)₂;

R⁷ is -CF₃ or R⁶;

R⁸ is hydrogen or X³-R⁹;

each R¹¹ is independently hydrogen or C₁₋₇ alkyl, or two R¹¹'s on same carbon atom are joined to form a cycloalkyl ring of 3 to 6 carbon atoms;

R¹² is hydrogen, C₁₋₇ alkyl or -CH₂R²¹;

R¹³ is C₁₋₇ alkyl or -(CH₂)₁R²¹;

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R^{14} is $-CF_3$ or R^{13} ;
 R^{15} is hydrogen, $-C(O)R^{16}$, R^{13} , or two R^{15} 's on the same nitrogen may be joined to form a monocyclic heterocyclic ring of 4 to 6 atoms containing up to 2 heteroatoms chosen from O, S or N;
 R^{16} is hydrogen, $-CF_3$, C_{1-7} alkyl, C_{2-7} alkenyl, C_{2-7} alkynyl or $-(CH_2)_sR^{21}$;
 R^{17} is $-(CH_2)_s-C(R^{18}R^{18})-(CH_2)_s-R^{19}$ or $-CH_2C(O)NR^{15}R^{15}$;
 R^{18} is hydrogen or C_{1-7} alkyl;
 R^{19} is a) a monocyclic or bicyclic heterocyclic ring containing from 3 to 9 nuclear carbon atoms and 1 or 2 nuclear hetero-atoms selected from N, S or O and with each ring in the heterocyclic radical being formed of 5 or 6 atoms, or b) the radical $W-R^{20}$;
 R^{20} is alkyl or $C(O)R^{23}$;
 R^{21} is phenyl substituted with 1 or 2 R^{22} groups;
 R^{22} is hydrogen, halogen, C_{1-7} alkyl, C_{1-7} alkoxy, C_{1-7} alkylthio, C_{1-7} alkylsulfonyl, C_{1-7} alkylcarbonyl, $-CF_3$, $-CN$, $-NO_2$ or $-N_3$;
 R^{23} is alkyl, cycloalkyl, monocyclic monoheterocyclic ring;
 R^{24} is the residual structure of a standard amino acid, or R^{18} and R^{24} attached to the same N can cyclize to form a proline residue;
m is 0 to 1;
n is 0 to 3;
p is 1 to 3 when m is 1;
p is 0 to 3 when m is 0;
r is 0 to 2;
s is 0 to 3;
t is 0 to 2;
u is 0 to 3;
v is 0 or 1;
W is O, S or NR^{15} ;
 X^1 is O, or NR^{15} ;
 X^2 is $C(O)$, $CR^{11}R^{11}$, S, $S(O)$ or $S(O)_2$;
 X^3 is $C(O)$, $CR^{11}R^{11}$, $S(O)_2$ or a bond;
Y is X^1 or X^2 ;
Q is $-CO_2R^{12}$, $-C(O)NHS(O)_2R^{14}$, $-NHS(O)_2R^{14}$, $-S(O)_2NHR^{15}$, $-C(O)NR^{15}R^{15}$, $-CO_2R^{17}$, $-C(O)NR^{18}R^{24}$, $-CH_2OH$, or 1H- or 2H-tetrazol-5-yl;
and the pharmaceutically acceptable salts thereof.

2. A compound of claim 1 wherein:

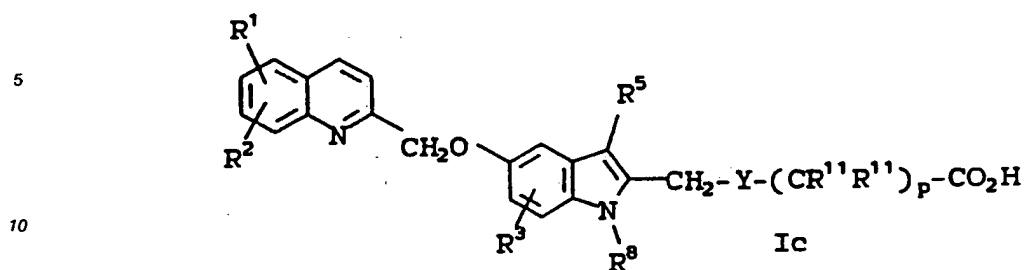
R^1 , R^2 , R^3 and R^4 are hydrogen;
 R^5 is X^2-R^7 or OR^6 ;
 R^7 is R^6 ;
 R^8 is R^9 ;
 R^{10} is hydrogen as halogen;
m is O;
n is 1 to 3;
u is O in R^6 and 1 in R^9 ;
v is O;
 X^2 is $CR^{11}R^{12}$ or S;
Q is $-CO_2R^{12}$;
or a pharmaceutically acceptable salt thereof.

3. A compound of claim 1 wherein n is o, m is o, v is o and Q is CO_2H or a pharmaceutically acceptable salt thereof.

4. A compound of any of Claims 1 to 3 wherein R^5 is SR^7 .

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5. The compound of Claim 1 of the formula Ic wherein the substituents are as follows:



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R ¹	R ²	R ³	R ⁵	R ⁸	Y-(CR ¹¹ R ¹¹) _p
6-Cl	7-Cl	H	Me	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	7-Cl	H	Me	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	4-allyl	-S-t-Bu	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	4-allyl	H	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	H	-O-i-Pr	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CH ₂ OCH(Me);
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CHMe;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-S-t-Bu	C(Me) ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CH ₂ OCH ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CHEt;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-Cl	CHMe;
H	H	H	-C(O)CH ₂ -t-Bu	H	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-CF ₃	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-3-OMe	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ CHCH ₂	CH ₂ OCH(Me);
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-OMe	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	Me	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-F	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-Br	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-I	C(Me) ₂ or
H	H	H	-t-Bu	-CH ₂ Ph-4-F	C(Me) ₂ ;

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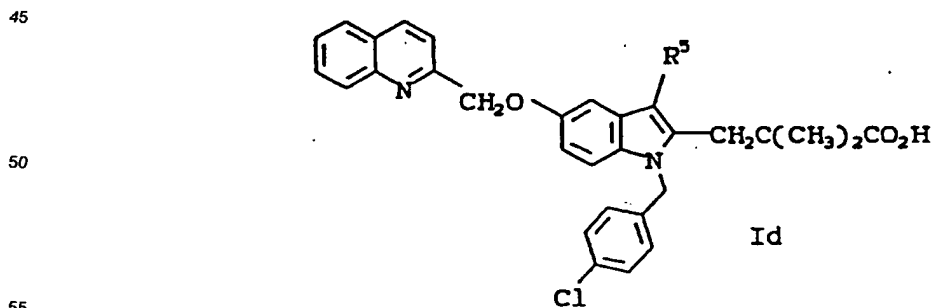
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and the pharmaceutically acceptable salts thereof.

6. The compound of Claim 1 of the Formula Id:



wherein R⁵ is as defined in Claim 1, are the pharmaceutically acceptable salts thereof.

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7. A compound of Claim 6 wherein the substituent R⁵ is: hydrogen, -Me, -S-t-Bu, -SPh, -S(O)₂Ph, -S(O)-Ph, C(O)Ph, -CH₂Ph, -C(O)CH₂-t-Bu, -O-i-Pr, -C(O)-CF₃, -S(O)₂-t-Bu, -S(O)-t-Bu, -C(O)Ph-4-t-Bu, -C(O)-Ph-4-Cl, t-Bu, -C(O)Me, -C(O)-c-Pr, -C(O)CH₂CH₂-c-C₅H₉, -C(O)CH₂CH(Me)₂, -C(O)Et, -C(O)CH(Me)₂, -C(O)C(Me)₃, -C(O)CH₂Ph, -C(Me)₂Pr, -C(Me)₂Et, -CH(Me)₂, -c-Pr, -(l-Me)-c-Pr, -c-C₅H₉, -c-C₆H₁₁, -C(Me)₂Ph, -C(Me)₂Ph-4-Cl, -1-Ad, -CH₂-1-Ad or -CH₂CH₂-t-Bu.
8. A compound according to Claim 1 which is:
- 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-quinolin-2-ylmethoxy]indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-methyl-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-t-butylthiobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-(phenylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-(phenylsulfonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethyl propanoic acid, N-oxide;
 - 3-[N-(p-chlorobenzyl)-3-(phenylsulfonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-(phenylsulfinyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-benzoyl-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-benzyl-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 2-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]ethoxyethanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-(3,3-dimethyl-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2-methylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-methyl-5-(6,7-dichloroquinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-methyl-5-(7-chloroquinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-4-allyl-5-(quinolin-2-ylmethoxy)-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-4-allyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-6-(quinolin-2-ylmethoxy)-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-4-(quinolin-2-ylmethoxy)-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-7-(quinolin-2-ylmethoxy)-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 2-[2-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]ethoxy]propanoic acid;
 - 3-[N-(p-chlorobenzyl)-4-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-methyl-3-(p-chlorobenzoyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-methyl-3-(p-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(4-chlorobenzyl)-3-i-propoxy-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(4-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-yl-methoxy)indol-2-yl]-2-ethylpropanoic acid;
 - 3-[N-(4-chlorobenzyl)-3-trifluoroacetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(4-chlorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2-methylpropanoic acid;
 - 3-[3-(3,3-dimethyl-1-oxo-1-butyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(4-trifluoromethylbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-benzyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(3-methoxybenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-allyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(4-methoxybenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;

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- 3-[N-methyl-3-(3,3-dimethyl-1-oxo-3-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid.
- 3-[N-(phenylsulfonyl)-3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 5 3-[N-benzyl-3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-(t-butylsulfonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-(t-butylsulfinyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 10 3-[N-allyl-3-(4-chlorobenzyl)-6-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(n-propyl)-3-(4-chlorobenzyl)-6-(quinoline-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-ethyl-3-(4-chlorobenzyl)-6-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-(4-t-butylbenzoyl)-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 15 3-[N-(4-chlorobenzyl)-3-(4-chlorobenzoyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-(1,1-dimethylethyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 20 3-[N-(4-chlorobenzyl)-3-acetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid
- 3-[N-(4-chlorobenzyl)-3-cyclopropanecarbonyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-(3-cyclopentylpropanoyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 25 3-[N-(4-chlorobenzyl)-3-(1-methylethyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-cyclopropyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-(1-methyl-1-cyclopropyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 30 3-[N-(4-chlorobenzyl)-3-cyclopentyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-cyclohexyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-(α,α -dimethylbenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-(2-{4-chloro- α,α -dimethylbenzyl})-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 35 3-[N-(4-chlorobenzyl)-3-(1-adamantyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-((1-adamantyl)methyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 40 3-[N-(1,1-dimethylethyl)-3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(1,1-dimethylpropyl)-3-(4-chlorobenzyl)-6-(quinoline-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-(3-methylbutanoyl)-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 45 3-[N-(4-chlorobenzyl)-3-propanoyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-(2-methylpropanoyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-trimethylacetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 50 3-[N-(4-chlorobenzyl)-3-phenylacetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-fluorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-bromobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 55 3-[N-(4-iodobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
- 3-[N-(4-chlorobenzyl)-3-(1,1-dimethylbutyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-

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dimethylpropanoic acid,

3-[N-(4-chlorobenzyl)-3-(1,1-dimethylpropyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-

dimethylpropanoic acid,

3-[N-(3-fluorobenzyl)-3-(1,1-dimethylethyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic

5 acid,

3-[N-(4-chlorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-

diethylpropanoic acid,

methyl 3-[N-(4-chlorobenzyl)-3,6-bis(acetyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2 dimethyl pro-

panoate or

10 methyl 3-[N-(4-chlorobenzyl)-3,6-bis (cyclopropanecarbonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-

dimethyl propanoate,

or a pharmaceutically acceptable salt thereof.

9. 3-[N-(p-Chlorobenzyl)-3-(t-butylthio)-5-quinolin-2-ylmethoxy)indole-2-yl]-2, 2-dimethylpropanoic acid or a
15 pharmaceutically acceptable salt thereof.

10. A pharmaceutically acceptable salt as claimed in any of claims 1 to 8 which is a sodium salt.

11. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 1
20 and a pharmaceutically acceptable carrier.

12. A pharmaceutical composition of Claim 11 additionally comprising an effective amount of a second
active ingredient selected from the group consisting of non-steroidal anti-inflammatory drugs; peripheral
analgesic agents; cyclooxygenase inhibitors; leukotriene antagonists; leukotriene biosynthesis inhibitors;
25 H₂-receptor antagonists; antihistaminic agents; prostaglandin antagonists; thromboxane antagonists;
thromboxane synthetase inhibitors; and ACE antagonists.

13. The use of a compound of any of claims 1 to 10 for the preparation of a medicament useful for
preventing the synthesis, the action, or the release of SRS-A or leukotrienes in a mammal.
30

14. The use of a compound of any of claims 1 to 10 for the preparation of a medicament useful for treating
asthma in a mammal.

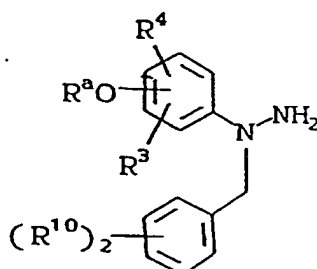
15. The use of a compound of any of claims 1 to 10 for the preparation of a medicament useful for treating
35 inflammatory diseases of the eye in a mammal.

16. The use of a compound of any of Claims 1 to 10 for the preparation of a medicament adapted for
administration of a compound of any of Claims 1 to 10 useful for treating inflammatory bowel disease in
a patient, treating arthritis in a patient, treating allergic rhinitis in a patient, treating allergic conjunctivitis
40 in a patient or treating renal insufficiency arising from ischaemia induced by immunological or chemical
toxicology in a patient.

17. A compound of the formula:

45

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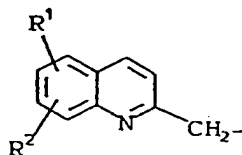
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wherein:

R^a is t-butylcarbonyl or

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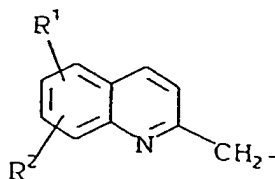


R¹, R², R³, R⁴ and R¹⁰ are as defined in Claim 1.

10

18. A compound according to Claim 17 wherein R^a is

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20

19. 1-(p-Chlorobenzyl)-1-[4-(quinolin-2-yl-methoxy)phenyl]hydrazine.

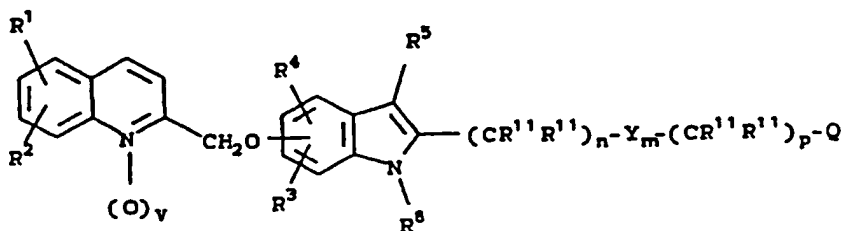
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Claims for the following Contracting States : ES, GR

1. A process for preparing a compound of the formula I:

30

35



40

I

wherein:

R¹, R², R³, R⁴ and R¹⁰ are independently hydrogen, halogen C₁₋₇ alkyl, C₂₋₇ alkenyl, C₂₋₇ alkynyl, -CF₃, -CN, -NO₂, -N₃, -C(OH)R¹¹R¹¹, -CO₂R¹², -SR¹⁴, -S(O)R¹⁴, -S(O)₂R¹⁴, -S(O)₂NR¹⁵R¹⁵, -OR¹⁵, -NR¹⁵R¹⁵, -C(O)R¹⁶ or -(CH₂)₁R²¹;

45

R⁵ is hydrogen, -CH₃, -CF₃, -C(O)H, X¹-R⁶ or X²-R⁷;

R⁶ and R⁹ are independently: alkyl, -(CH₂)_uPh(R¹⁰)₂ or -(CH₂)_uTh(R¹⁰)₂;

R⁷ is -CF₃ or R⁶;

R⁸ is hydrogen or X³-R⁹;

50

each R¹¹ is independently hydrogen or C₁₋₇ alkyl, or two R¹¹'s on same carbon atom are joined to form a cycloalkyl ring of 3 to 6 carbon atoms;

R¹² is hydrogen, C₁₋₇ alkyl or -CH₂R²¹;

R¹³ is C₁₋₇ alkyl or -(CH₂)₁R²¹;

R¹⁴ is -CF₃ or R¹³;

55

R¹⁵ is hydrogen, -C(O)R¹⁶, R¹³, or two R¹⁵'s on the same nitrogen may be joined to form a monocyclic heterocyclic ring of 4 to 6 atoms containing up to 2 heteroatoms chosen from O, S or N;

R¹⁶ is hydrogen, -CF₃, C₁₋₇ alkyl, C₂₋₇ alkenyl, lower alkynyl or -(CH₂)₁R²¹;

R¹⁷ is -(CH₂)_s-C(R¹⁸R¹⁸)-(CH₂)_s-R¹⁹ or -CH₂C(O)NR¹⁵R¹⁵;

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R¹⁸ is hydrogen or C₁₋₇ alkyl;

R¹⁹ is a) a monocyclic or bicyclic heterocyclic ring containing from 3 to 9 nuclear carbon atoms and 1 or 2 nuclear hetero-atoms selected from N, S or O and with each ring in the heterocyclic radical being formed of 5 or 6 atoms, or b) the radical W-R²⁰;

5 R²⁰ is alkyl or C(O)R²³;

R²¹ is phenyl substituted with 1 or 2 R²² groups;

R²² is hydrogen, halogen, C₁₋₇ alkyl, C₁₋₇ alkoxy, C₁₋₇ alkylthio, C₁₋₇ alkylsulfonyl, C₁₋₇ alkylcarbonyl, -CF₃, -CN, -NO₂ or -N₃;

R²³ is alkyl, cycloalkyl, monocyclic monoheterocyclic ring;

10 R²⁴ is the residual structure of a standard amino acid, or R¹⁸ and R²⁴ attached to the same N can cyclize to form a proline residue;

m is 0 to 1;

n is 0 to 3;

p is 1 to 3 when m is 1;

15 p is 0 to 3 when m is 0;

r is 0 to 2;

s is 0 to 3;

t is 0 to 2;

u is 0 to 3;

20 v is 0 or 1;

W is O, S or NR¹⁵;

X¹ is O, or NR¹⁵;

X² is C(O), CR¹¹R¹¹, S, S(O) or S(O)₂;

X³ is C(O), CR¹¹R¹¹, S(O)₂ or a bond;

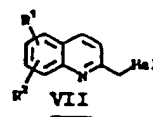
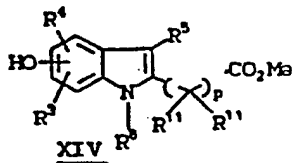
25 Y is X¹ or X²;

Q is -CO₂R¹², -C(O)NHS(O)₂R¹⁴, -NHS(O)₂R¹⁴, -S(O)₂NHR¹⁵, -C(O)NR¹⁵R¹⁵, -CO₂R¹⁷, -C(O)NR¹⁸R²⁴, -CH₂OH, or 1H- or 2H-tetrazol-5-yl;

and the pharmaceutically acceptable salts thereof,

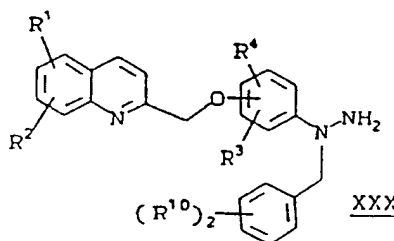
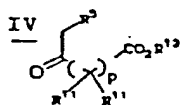
which comprises:

30 coupling a compound of formula XIV with a compound of formula VII



40 to yield a compound of the formula (I) wherein (CR¹¹R¹¹)_n-Y_m-(CR¹¹R¹¹)_p-Q is (CR¹¹R¹¹)_pCO₂Me and thereafter if desired transforming the group (CR¹¹R¹¹)_pCO₂Me into another group of the formula (CR¹¹R¹¹)_n-Y_m-(CR¹¹R¹¹)_p-Q.

45 2. A process for preparing the compound of formula I wherein V is O and R⁸ is CH₂C₆H₃(R¹⁰)₂, as defined in Claim 1, which comprises reacting the compound of formula XXX with the ketone of formula IV by a Fischer reaction, to provide the compound of formula I:

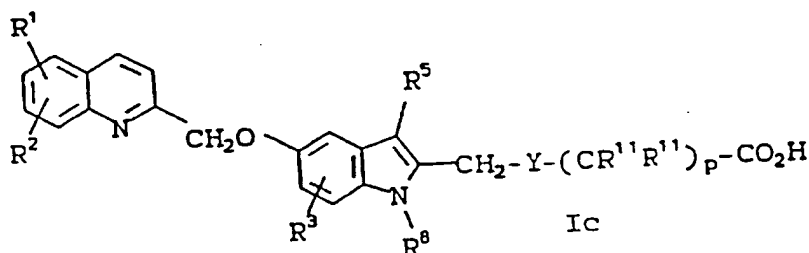


wherein V is O and R⁸ is CH₂C₆H₃(R¹⁰)₂ and (CR¹¹R¹¹)_n-Y_m-(CR¹¹R¹¹)_p-Q is (CR¹¹R¹¹)_pCO₂Me and

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thereafter if desired transforming the group $(CR^{11}R^{11})_pCO_2Me$ into another group of the formula $(CR^{11}R^{11})_n-Y_m-(CR^{11}R^{11})_p-Q$.

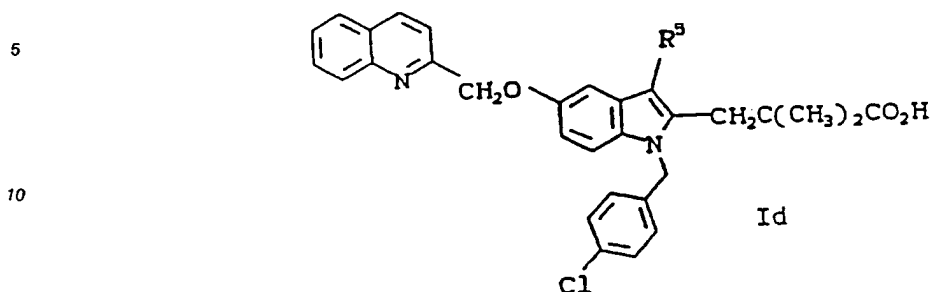
3. A process as claimed in Claim 1 for preparing a compound of the formula (I) as defined in Claim 1 wherein n is 0, m is 0 and Q is CO_2H or a pharmaceutically acceptable salt thereof.
4. A process as claimed in Claim 2 for preparing a compound of the formula (I) as defined in Claim 2 wherein n is 0, m is 0 and Q is CO_2H or a pharmaceutically acceptable salt thereof.
5. A process according to any of Claims 1 to 4 for preparing the compound of the formula Ic or a pharmaceutically acceptable salt thereof wherein the substituents are as follows:



R ¹	R ²	R ³	R ⁵	R ⁸	Y-(CR ¹¹ R ¹¹) _p
6-Cl	7-Cl	H	Me	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	7-Cl	H	Me	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	4-allyl	-S-t-Bu	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	4-allyl	H	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	H	-O-i-Pr	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CH ₂ OCH(Me);
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CHMe;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-S-t-Bu	C(Me) ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CH ₂ OCH ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CHEt;
H	H	H	-C(O)CH ₂ -t-	-CH ₂ Ph-4-Cl	CHMe;
H	H	H	-C(O)CH ₂ -t-Bu	H	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-CF ₃	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-3-OMe	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ CHCH ₂	CH ₂ OCH(Me);
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-OMe	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	Me	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-F	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-Br	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-I	C(Me) ₂ or
H	H	H	-t-Bu	-CH ₂ Ph-4-F	C(Me) ₂ .

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6. A process according to any of Claims 1 to 4 for preparing a compound of the formula Id:



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or a pharmaceutically acceptable salt thereof wherein R⁵ is as defined in Claim 1.

7. A process according to Claim 6 for preparing the compound of the formula Id wherein the substituent R⁵ is: hydrogen, -Me, -S-t-Bu, -SPh, -S(O)₂Ph, -S(O)Ph, C(O)Ph, -CH₂Ph, -C(O)CH₂-t-Bu, -O-i-Pr, -C(O)-CF₃, -S(O)₂-t-Bu, -S(O)-t-Bu, -C(O)Ph-4-t-Bu, -C(O)Ph-4-Cl, t-Bu, -C(O)Me, -C(O)-c-Pr, -C(O)-CH₂CH₂-c-C₅H₉, -C(O)CH₂CH(Me)₂, -C(O)Et, -C(O)CH(Me)₂, -C(O)C(Me)₃, -C(O)CH₂Ph, -C(Me)₂Pr, -C(Me)₂Et, -CH(Me)₂, -c-Pr, -(l-Me)-c-Pr, -c-C₅H₉, -c-C₆H₁₁, -C(Me)₂Ph, -C(Me)₂Ph-4-Cl, -1-Ad, -CH₂-1-Ad or -CH₂CH₂-t-Bu.
- 20

- 25 8. A process according to any of Claims 1 to 4 for preparing a compound which is:
- 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-methyl-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-t-butylthiobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-(phenylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-(phenylsulfonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethyl propanoic acid, N-oxide;
 - 3-[N-(p-chlorobenzyl)-3-(phenylsulfonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-(phenylsulfinyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-benzoyl-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-benzyl-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 2-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]ethoxyethanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-(3,3-dimethyl-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2-methylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-methyl-5-(6,7-dichloroquinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-3-methyl-5-(7-chloroquinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-4-allyl-5-(quinolin-2-yl-methoxy)-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-4-allyl-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-6-(quinolin-2-ylmethoxy)-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-4-(quinolin-2-ylmethoxy)-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-(p-chlorobenzyl)-7-(quinolin-2-ylmethoxy)-3-(t-butylthio)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 2-[2-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]ethoxy]propanoic acid;
 - 3-[N-(p-chlorobenzyl)-4-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid;
 - 3-[N-methyl-3-(p-chlorobenzoyl)-6-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
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3-[N-methyl-3-(p-chlorobenzyl)-6-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
3-[N-(4-chlorobenzyl)-3-i-propoxy-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
3-[N-(4-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-yl-methoxy)indol-2-yl]-2-ethylpropanoic acid,
3-[N-(4-chlorobenzyl)-3-trifluoroacetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic
5 acid,
3-[N-(4-chlorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2-
methylpropanoic acid,
3-[3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
3-[N-(4-trifluoromethylbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-
10 dimethylpropanoic acid,
3-[N-benzyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic
acid,
3-[N-(3-methoxybenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-
dimethylpropanoic acid,
15 3-[N-allyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic
acid,
3-[N-(4-methoxybenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-
dimethylpropanoic acid,
3-[N-methyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic
20 acid,
3-[3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid.
3-[N-(phenylsulfonyl)-3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic
acid,
3-[N-benzyl-3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
25 3-[N-(4-chlorobenzyl)-3-(t-butylsulfonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic
acid,
3-[N-(4-chlorobenzyl)-3-(t-butylsulfinyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic
acid,
3-[N-allyl-3-(4-chlorobenzyl)-6-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
30 3-[N-(n-propyl)-3-(4-chlorobenzyl)-6-(quinoline-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
3-[N-ethyl-3-(4-chlorobenzyl)-6-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
3-[N-(4-chlorobenzyl)-3-(4-t-butylbenzoyl)-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-dimethylpropanoic
acid,
3-[N-(4-chlorobenzyl)-3-(4-chlorobenzoyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic
35 acid,
3-[N-(4-chlorobenzyl)-3-(1,1-dimethylethyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-
dimethylpropanoic acid,
3-[N-(4-chlorobenzyl)-3-acetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid
3-[N-(4-chlorobenzyl)-3-cyclopropanecarbonyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-
40 dimethylpropanoic acid,
3-[N-(4-chlorobenzyl)-3-(3-cyclopentylpropanoyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-
dimethylpropanoic acid,
3-[N-(4-chlorobenzyl)-3-(3-methylbutanoyl)-5-(quinolin-2-yl-methoxy)indol-2-yl]-2,2-
dimethylpropanoic acid,
45 3-[N-(4-chlorobenzyl)-3-propanoyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
3-[N-(4-chlorobenzyl)-3-(2-methylpropanoyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-
dimethylpropanoic acid,
3-[N-(4-chlorobenzyl)-3-trimethylacetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic
acid,
50 3-[N-(4-chlorobenzyl)-3-phenylacetyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
3-[N-(4-fluorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-
dimethylpropanoic acid,
3-[N-(4-bromobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-
dimethylpropanoic acid,
55 3-[N-(4-iodobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-
dimethylpropanoic acid,
3-[N-(4-chlorobenzyl)-3-(1,1-dimethylbutyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-
dimethylpropanoic acid,

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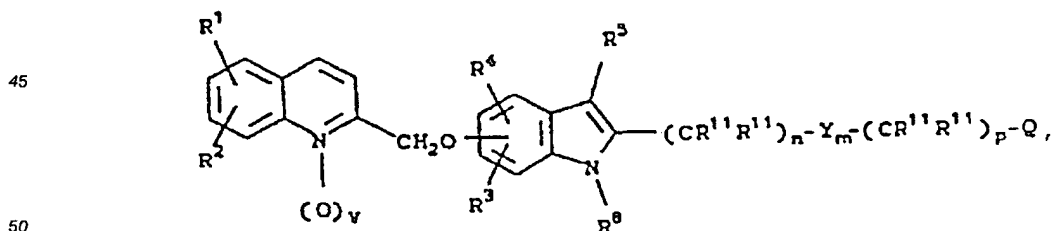
- 3-[N-(4-chlorobenzyl)-3-(1,1-dimethylpropyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
 3-[N-(3-fluorobenzyl)-3-(1,1-dimethylethyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
 5 3-[N-(4-chlorobenzyl)-3-(1-methylethyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
 3-[N-(4-chlorobenzyl)-3-cyclopropyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
 3-[N-(4-chlorobenzyl)-3-(1-methyl-1-cyclopropyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
 10 3-[N-(4-chlorobenzyl)-3-cyclopentyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
 3-[N-(4-chlorobenzyl)-3-cyclohexyl-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
 3-[N-(4-chlorobenzyl)-3-(α,α -dimethylbenzyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
 3-[N-(4-chlorobenzyl)-3-(2-{4-chloro- α,α -dimethylbenzyl})-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
 15 3-[N-(4-chlorobenzyl)-3-(1-adamantyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
 3-[N-(4-chlorobenzyl)-3-((1-adamantyl)methyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
 20 3-[N-(1,1-dimethylethyl)-3-(4-chlorobenzyl)-6-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
 3-[N-(1,1-dimethylpropyl)-3-(4-chlorobenzyl)-6-(quinoline-2-ylmethoxy)indol-2-yl]-2,2-dimethylpropanoic acid,
 3-[N-(4-chlorobenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-diethylpropanoic acid,
 25 methyl 3-[N-(4-chlorobenzyl)-3,6-bis(acetyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2 dimethyl propanoate or
 methyl 3-[N-(4-chlorobenzyl)-3,6-bis (cyclopropanecarbonyl)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2,2-dimethyl propanoate
 30 or a pharmaceutically acceptable salt thereof.

9. A process for the preparation of 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy) indol-2-yl]-2,2-dimethylpropanoic acid or a pharmaceutically acceptable salt thereof which comprises the hydrolysis of methyl 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinolin-2-ylmethoxy)indol-2-yl]-2, 2-dimethylpropanoate.
 35

Patentansprüche

Patentansprüche für folgende Vertragsstaaten : AT, BE, CH, DE, DK, FR, GB, IT, LI, LU, NL, SE

- 40 1. Verbindung der Formel I:



I

- 55 worin

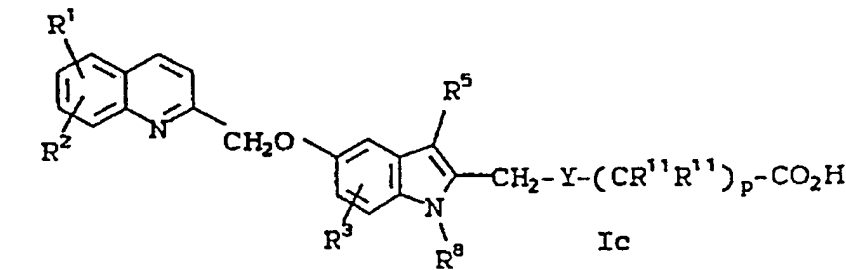
R¹, R², R³, R⁴ und R¹⁰ unabhängig für Wasserstoff, Halogen, C₁₋₇-Alkyl, C₂₋₇-Alkenyl, C₂₋₇-Alkynyl, -CF₃, -CN, -NO₂, -N₃, -C(OH)R¹¹R¹¹, -CO₂R¹², -SR¹⁴, -S(O)R¹⁴, -S(O)₂R¹⁴, -S(O)₂NR¹⁵R¹⁵, -OR¹⁵, -NR¹⁵R¹⁵, -C(O)R¹⁶ oder -(CH₂)₁R²¹ stehen;

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- R^5 für Wasserstoff, $-CH_3$, $-CF_3$, $-C(O)H$, X^1-R^6 oder X^2-R^7 steht;
 R^6 und R^9 unabhängig für Alkyl, $-(CH_2)_uPh(R^{10})_2$ oder $-(CH_2)_uTh(R^{10})_2$ stehen;
 R_7 für $-CF_3$ oder R^6 steht;
 R^8 für Wasserstoff oder X^3-R^9 steht;
 5 jedes R^{11} unabhängig für Wasserstoff oder C_1-7 -Alkyl steht, oder zwei R^{11} an demselben Kohlenstoff unter Bildung eines Cycloalkylrings mit 3 bis 6 Kohlenstoffatomen verknüpft sind;
 R^{12} für Wasserstoff, C_1-7 -Alkyl oder $-CH_2R^{21}$ steht;
 R^{13} für C_1-7 -Alkyl oder $-(CH_2)_rR^{21}$ steht;
 R^{14} für $-CF_3$ oder R^{13} steht;
 10 R^{15} für Wasserstoff, $-C(O)R^{16}$, R^{13} steht, oder zwei R^{15} an demselben Stickstoffatom unter Bildung eines monocyclischen heterocyclischen Rings mit 4 bis 6 Atomen, der bis zu zwei Heteroatome, ausgewählt aus O, S oder N enthält, verknüpft sein können;
 R^{16} für Wasserstoff, $-CF_3$, C_1-7 -Alkyl, C_2-7 -Alkenyl, C_2-7 -Alkynyl oder $-(CH_2)_rR^{21}$ steht;
 R^{17} für $-(CH_2)_s-C(R^{18}R^{18})-(CH_2)_s-R^{19}$ oder $-CH_2C(O)NR^{15}R^{15}$ steht;
 15 R^{18} für Wasserstoff oder C_1-7 -Alkyl steht;
 R^{19} für a) einen monocyclischen oder bicyclischen heterocyclischen Ring steht, der 3 bis 9 Ringkohlenstoffatome und 1 oder 2 Ringheteroatome, ausgewählt aus N, S oder O, enthält, und wobei jeder Ring in dem heterocyclischen Rest aus 5 oder 6 Atomen gebildet wird, oder b) den Rest $W-R^{20}$ steht;
 R^{20} für Alkyl oder $C(O)R^{23}$ steht;
 20 R^{21} für Phenyl, substituiert mit 1 oder 2 R^{22} -Gruppen, steht;
 R^{22} für Wasserstoff, Halogen, C_1-7 -Alkyl, C_1-7 -Alkoxy, C_1-7 -Alkylthio, C_1-7 -Alkylsulfonyl, C_1-7 -Alkylcarbonyl, $-CF_3$, $-CN$, $-NO_2$ oder $-N_3$ steht;
 R^{23} für Alkyl, Cycloalkyl, einen monocyclischen monoheterocyclischen Ring steht;
 R^{24} die Reststruktur einer Standardamino-säure darstellt, oder R^{18} und R^{24} , die an dasselbe N
 25 angeknüpft sind, unter Bildung eines Prolinrestes cyclisieren können;
 m für 0 bis 1 steht;
 n für 0 bis 3 steht;
 p für 1 bis 3 steht, wenn m für 1 steht;
 p für 0 bis 3 steht, wenn m für 0 steht;
 30 r für 0 bis 2 steht;
 s für 0 bis 3 steht;
 t für 0 bis 2 steht;
 u für 0 bis 3 steht;
 v für 0 bis 1 steht;
 35 W für O, S oder NR^{15} steht;
 X^1 für 0 oder NR^{15} steht;
 X^2 für $C(O)$, $CR^{11}R^{11}$, S, $S(O)$ oder $S(O)_2$ steht;
 X^3 für $C(O)$, $CR^{11}R^{11}$, $S(O)_2$ oder eine Bindung steht;
 Y für X^1 oder X^2 steht;
 40 Q für $-CO_2R^{12}$, $-C(O)NHS(O)_2R^{14}$, $-NHS(O)_2R^{14}$, $-S(O)_2NHR^{15}$, $-C(O)NR^{15}R^{15}$, $-CO_2R^{17}$, $-C(O)NR^{18}R^{24}$,
 $-CH_2OH$ oder 1H- oder 2H-Tetrazol-5-yl steht;
 und die pharmazeutisch verträglichen Salze davon.
2. Verbindung nach Anspruch 1, worin:
 45 R^1 , R^2 , R^3 und R^4 Wasserstoff bedeuten;
 R^5 X^2-R^7 oder OR^6 bedeutet;
 R^7 R^6 bedeutet;
 R^8 R^9 bedeutet;
 R^{10} Wasserstoff ebenso wie Halogen bedeutet;
 50 m für 0 steht;
 n eine Zahl von 1 bis 3 bedeutet;
 u für 0 in R^6 steht und für 1 in R^9 steht;
 v für 0 steht;
 X^2 $CR^{11}R^{12}$ oder S bedeutet;
 55 Q $-CO_2R^{12}$ bedeutet;
 oder ein pharmazeutisch verträgliches Salz davon.

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3. Verbindung nach Anspruch 1, worin n 0 bedeutet, m 0 bedeutet, v 0 bedeutet, und Q CO₂H bedeutet, oder ein pharmazeutisch verträgliches Salz davon.
4. Verbindung nach einem der Ansprüche 1 bis 3, worin R⁵ SR⁷ bedeutet.
5. Verbindung nach Anspruch 1 der Formel Ic, worin die Substituenten wie folgt lauten:

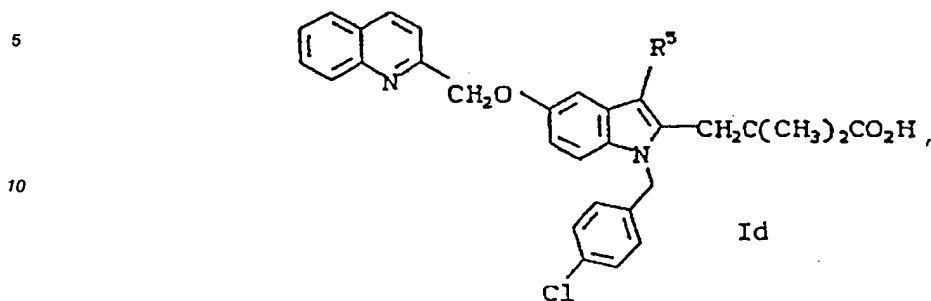


R ¹	R ²	R ³	R ⁵	R ⁸	Y-(CR ¹¹ R ¹¹) _p
6-Cl	7-Cl	H	Me	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	7-Cl	H	Me	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	4-Allyl	-S-t-Bu	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	4-Allyl	H	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	H	-O-i-Pr	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CH ₂ OCH(Me);
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CHMe;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-S-t-Bu	C(Me) ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CH ₂ OCH ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CHEt;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-Cl	CHMe;
H	H	H	-C(O)CH ₂ -t-Bu	H	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-CF ₃	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-3-OMe	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ CHCH ₂	CH ₂ OCH(Me);
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-OMe	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	Me	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-F	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-Br	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-I	C(Me) ₂ oder
H	H	H	-t-Bu	-CH ₂ Ph-4-F	C(Me) ₂ ;

und die pharmazeutisch verträglichen Salze davon.

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6. Verbindung nach Anspruch 1 der Formel Id:



worin R⁵ die in Anspruch 1 gegebene Definition besitzt, und die pharmazeutisch verträglichen Salze davon.

20 7. Verbindung nach Anspruch 6, worin der Substituent R⁵ für folgendes steht:

Wasserstoff, -Me, -S-t-Bu, -SPh, -S(O)₂Ph, -S(O)Ph, C(O)Ph, -CH₂Ph, -C(O)CH₂-t-Bu, -O-i-Pr, -C(O)-CF₃, -S(O)₂-t-Bu, S(O)-t-Bu, -C(O)Ph-4-t-Bu, -C(O)Ph-4-Cl, t-Bu, -C(O)Me, -C(O)-c-Pr, -C(O)-CH₂CH₂-c-C₅H₉, -C(O)CH₂CH(Me)₂, -C(O)Et, -C(O)CH(Me)₂, -C(O)C(Me)₃, -C(O)CH₂Ph, -C(Me)₂Pr, -C(Me)₂Et, -CH(Me)₂, -c-Pr, -(l-Me)-c-Pr, -c-C₅H₉, -c-C₆H₁₁, -C(Me)₂Ph, -C(Me)₂Ph-4-Cl, -1-Ad, -CH₂-1-Ad oder -CH₂CH₂-t-Bu.

25

8. Verbindung nach Anspruch 1, die folgendermaßen heißt:

3-[N-(p-Chlorbenzyl)-3-(t-butylthio)-5-chinolin-2-yl-methoxy]-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-methyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 30 3-[N-(p-t-Butylthiobenzyl)-3-(t-butylthio)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-(phenylthio)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-(phenylsulfonyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure-N-Oxid,
 35 3-[N-(p-Chlorbenzyl)-3-(Phenylsulfonyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-(phenylsulfinyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 40 3-[N-(p-Chlorbenzyl)-3-benzoyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-benzyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 2-[N-(p-Chlorbenzyl)-3-(t-butylthio)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-ethoxyethansäure,
 45 3-[N-(p-Chlorbenzyl)-3-(3,3-dimethyl-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-(t-butylthio)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2-methylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-methyl-5-(6,7-dichlorchinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 50 3-[N-(p-Chlorbenzyl)-3-methyl-5-(7-chlorchinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-4-allyl-5-(chinolin-2-yl-methoxy)-3-(t-butylthio)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-4-allyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-6-(chinolin-2-yl-methoxy)-3-(t-butylthio)-indol-2-yl]-2,2-dimethylpropansäure,
 55 3-[N-(p-Chlorbenzyl)-4-(chinolin-2-yl-methoxy)-3-(t-butylthio)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-7-(chinolin-2-yl-methoxy)-3-(t-butylthio)-indol-2-yl]-2,2-dimethylpropansäure,
 2-[2-[N-(p-Chlorbenzyl)-3-(t-butylthio)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-ethoxy]-propansäure,
 3-[N-(p-Chlorbenzyl)-4-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,

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- 3-[N-Methyl-3-(p-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-Methyl-3-(p-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Chlorbenzyl)-3-i-propoxy-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Chlorbenzyl)-3-(t-butylthio)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2-ethylpropansäure,
 5 3-[N-(4-Chlorbenzyl)-3-trifluoracetyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Chlorbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2-methylpropansäure,
 3-[3-(3,3-Dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Trifluormethylbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 10 3-[N-Benzyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(3-Methoxybenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-Allyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 15 3-[N-(4-Methoxybenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-Methyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 20 3-[3-(4-Chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(Phenylsulfonyl)-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-Benzyl-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 25 3-[N-(4-Chlorbenzyl)-3-(t-butylsulfonyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Chlorbenzyl)-3-(t-butylsulfinyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-Allyl-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 30 3-[N-(n-Propyl)-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-Ethyl-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Chlorbenzyl)-3-(4-t-butylbenzoyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Chlorbenzyl)-3-(4-chlorbenzoyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 35 3-[N-(4-Chlorbenzyl)-3-(1,1-dimethylethyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Chlorbenzyl)-3-acetyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Chlorbenzyl)-3-cyclopropanecarbonyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 40 3-[N-(4-Chlorbenzyl)-3-(3-cyclopentylpropanoyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Chlorbenzyl)-3-(1-methylethyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure
 3-[N-(4-Chlorbenzyl)-3-cyclopropyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure
 45 3-[N-(4-Chlorbenzyl)-3-(1-methyl-1-cyclopropyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure
 3-[N-(4-Chlorbenzyl)-3-cyclopentyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure
 3-[N-(4-Chlorbenzyl)-3-cyclohexyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure
 3-[N-(4-Chlorbenzyl)-3-(α,α -dimethylbenzyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure
 50 3-[N-(4-Chlorbenzyl)-3-(2-{4-chlor- α,α -dimethylbenzyl})-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure
 3-[N-(4-Chlorbenzyl)-3-(1-adamantyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure
 3-[N-(4-Chlorbenzyl)-3-((1-adamantyl)-methyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure
 55 3-[N-(1,1-Dimethylethyl)-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure
 3-[N-(1,1-Dimethylpropyl)-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure

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dimethylpropansäure

3-[N-(4-Chlorbenzyl)-3-(3-methylbutanoyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

dimethylpropansäure,

3-[N-(4-Chlorbenzyl)-3-propanoyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,

5 3-[N-(4-Chlorbenzyl)-3-(2-methylpropanoyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

dimethylpropansäure,

3-[N-(4-Chlorbenzyl)-3-trimethylacetyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,

3-[N-(4-Chlorbenzyl)-3-phenylacetyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,

3-[N-(4-Fluorbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

10 dimethylpropansäure,

3-[N-(4-Brombenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

dimethylpropansäure,

3-[N-(4-Iodbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

dimethylpropansäure,

15 3-[N-(4-Chlorbenzyl)-3-(1,1-dimethylbutyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

dimethylpropansäure,

3-[N-(4-Chlorbenzyl)-3-(1,1-dimethylpropyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

dimethylpropansäure,

3-[N-(3-Fluorbenzyl)-3-(1,1-dimethylethyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

20 dimethylpropansäure,

3-[N-(4-Chlorbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

diethylpropansäure,

Methyl-3-[N-(4-chlorbenzyl)-3,6-bis(acetyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

dimethylpropanoat oder

25 Methyl-3-[N-(4-chlorbenzyl)-3,6-bis(cyclopropanecarbonyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

dimethylpropanoat,

oder ein pharmazeutisch verträgliches Salz davon.

9. 3-[N-(p-Chlorbenzyl)-3-(t-butylthio)-5-chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure oder
30 ein pharmazeutisch verträgliches Salz davon.

10. Pharmazeutisch verträgliches Salz nach einem der Ansprüche 1 bis 8, das das Natriumsalz ist.

11. Pharmazeutische Zusammensetzung, umfassend eine therapeutisch wirksame Menge einer Verbindung
35 nach Anspruch 1 und einen pharmazeutisch verträglichen Trägerstoff.

12. Pharmazeutische Zusammensetzung nach Anspruch 11, die außerdem eine wirksame Menge eines
40 zweiten aktiven Bestandteils umfaßt, ausgewählt aus der Gruppe, bestehend aus nichtsteroiden antiinflammatorischen Wirkstoffen, peripher wirksamen Analgetika, Cyclooxygenase-Hemmern, Leukotrien-Anatagonisten, Leukotrien-Biosynthesehemmern, H₂-Rezeptorantagonisten, Antihistaminika, Prostaglandin-Antagonisten, Thromboxan-Antagonisten, Thromboxan-Synthetasehemmern und ACE-Antagonisten.

13. Verwendung einer Verbindung nach einem der Ansprüche 1 bis 10 zur Herstellung eines Medikaments,
45 das zur Prävention von Synthese, Wirkung oder Freisetzung von SRS-A oder Leukotrienen bei einem Säuger geeignet ist.

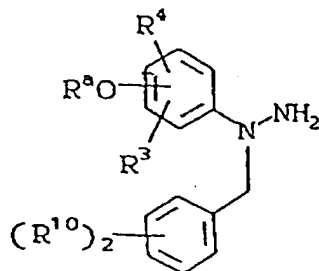
14. Verwendung einer Verbindung nach einem der Ansprüche 1 bis 10 zur Herstellung eines Medikaments,
das zur Behandlung von Asthma bei einem Säuger geeignet ist.

50 15. Verwendung einer Verbindung nach einem der Ansprüche 1 bis 10 zur Herstellung eines Medikaments,
das zur Behandlung entzündlicher Augenkrankheiten bei einem Säuger geeignet ist.

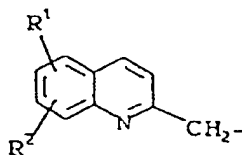
16. Verwendung einer Verbindung nach einem der Ansprüche 1 bis 10 zur Herstellung eines Medikaments,
55 das zur Verabreichung einer Verbindung nach einem der Ansprüche 1 bis 10 angepaßt ist, das geeignet ist zur Behandlung entzündlicher Darmerkrankung bei einem Patienten, zur Behandlung von Arthritis bei einem Patienten, zur Behandlung allergischer Rhinitis bei einem Patienten, zur Behandlung allergischer Konjunktivitis bei einem Patienten oder zur Behandlung einer Niereninsuffizienz bei einem Patienten aufgrund von Ischaemie, ausgelöst durch immunologische oder chemische Vergiftung.

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17. Verbindung der Formel:

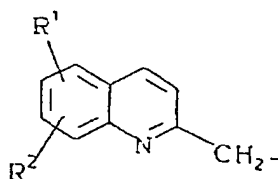


worin:
R^a für t-Butylcarbonyl oder



steht;
R¹, R², R³, R⁴ und R¹⁰ die in Anspruch 1 gegebenen Definitionen besitzen.

18. Verbindung nach Anspruch 17, worin R^a für



steht.

19. 1-(p-Chlorbenzyl)-1-[4-chinolin-2-ylmethoxy)-phenyl]-hydrazin.

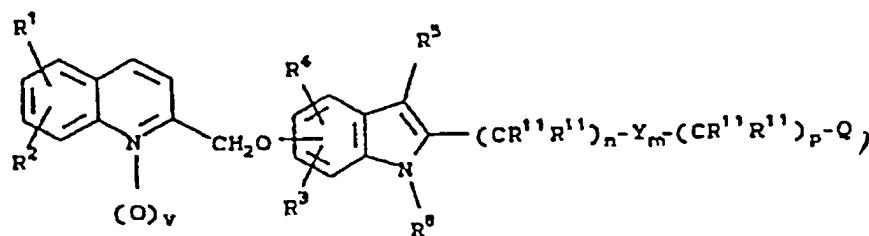
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Patentansprüche für folgende Vertragsstaaten : ES, GR

1. Verfahren zur Herstellung einer Verbindung der Formel I:

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I

- worin R¹, R², R³, R⁴ und R¹⁰ unabhängig für Wasserstoff, Halogen, C₁-7-Alkyl, C₂-7-Alkenyl, C₂-7-Alkynyl, -CF₃, -CN, -NO₂, -N₃, -C(OH)R¹¹, -CO₂R¹², -SR¹⁴, -S(O)R¹⁴, -S(O)₂R¹⁴, -S(O)₂NR¹⁵R¹⁵,
 -OR¹⁵, -NR¹⁵R¹⁵, -C(O)R¹⁶ oder -(CH₂)_tR²¹ stehen;
 R⁵ für Wasserstoff, -CH₃, -CF₃, -C(O)H, X¹-R⁶ oder X²-R⁷ steht;
 R⁶ und R⁹ unabhängig für Alkyl, -(CH₂)_uPh(R¹⁰)₂ oder -(CH₂)_uTh(R¹⁰)₂ stehen;
 R₇ für -CF₃ oder R⁶ steht;
 R⁸ für Wasserstoff oder X³-R⁹ steht;
 jedes R¹¹ unabhängig für Wasserstoff oder C₁-7-Alkyl steht, oder zwei R¹¹ mit demselben Kohlenstoff unter Bildung eines Cycloalkylrings mit 3 bis 6 Kohlenstoffatomen verknüpft sind;
 R¹² für Wasserstoff, C₁-7-Alkyl oder -CH₂R²¹ steht;
 R¹³ für C₁-7-Alkyl oder -(CH₂)_rR²¹ steht;
 R¹⁴ für -CF₃ oder R¹³ steht;
 R¹⁵ für Wasserstoff, -C(O)R¹⁶, R¹³ steht, oder zwei R¹⁵ mit demselben Stickstoffatom unter Bildung eines monocyclischen heterocyclischen Rings von 4 bis 6 Atomen, der bis zu zwei Heteroatome, ausgewählt aus O, S oder N enthält, verknüpft sein können;
 R¹⁶ für Wasserstoff, -CF₃, C₁-7-Alkyl, C₂-7-Alkenyl, Niedrigalkynyl oder -(CH₂)_sR²¹ steht;
 R¹⁷ für -(CH₂)_s-C(R¹⁸R¹⁸)-(CH₂)_s-R¹⁹ oder -CH₂C(O)NR¹⁵R¹⁵ steht;
 R¹⁸ für Wasserstoff oder C₁-7-Alkyl steht;
 R¹⁹ für a) einen monocyclischen oder bicyclischen heterocyclischen Ring steht, der 3 bis 9 Ringkohlenstoffe und 1 oder 2 Ringheteroatome, ausgewählt aus N, S oder O, enthält, und wobei jeder Ring in dem heterocyclischen Rest aus 5 oder 6 Atomen gebildet wird, oder b) den Rest W-R²⁰ steht;
 R²⁰ für Alkyl oder C(O)R²³ steht;
 R²¹ für Phenyl, substituiert mit 1 oder 2 R²²-Gruppen, steht;
 R²² für Wasserstoff, Halogen, C₁-7-Alkyl, C₁-7-Alkoxy, C₁-7-Alkylthio, C₁-7-Alkylsulfonyl, C₁-7-Alkyl-carbonyl, -CF₃, -CN, -NO₂ oder -N₃ steht;
 R²³ für Alkyl, Cycloalkyl, einen monocyclischen monoheterocyclischen Ring steht;
 R²⁴ die Reststruktur einer Standard-Aminosäure darstellt, oder R¹⁸ und R²⁴, die an dasselbe N angeknüpft sind, unter Bildung eines Prolinrestes cyclisieren können;
 m für 0 bis 1 steht;
 n für 0 bis 3 steht;
 p für 1 bis 3 steht, wenn m für 1 steht;
 p für 0 bis 3 steht, wenn m für 0 steht;
 r für 0 bis 2 steht;
 s für 0 bis 3 steht;
 t für 0 bis 2 steht;
 u für 0 bis 3 steht;
 v für 0 bis 1 steht;
 W für O, S oder NR¹⁵ steht;
 X¹ für O oder NR¹⁵ steht;
 X² für C(O), CR¹¹R¹¹, S, S(O) oder S(O)₂ steht;
 X³ für C(O), CR¹¹R¹¹, S(O)₂ oder eine Bindung steht;

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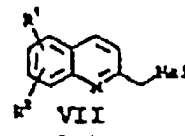
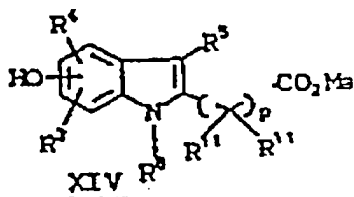
Y für X¹ oder X² steht;

Q für -CO₂R¹², -C(O)NHS(O)₂R¹⁴, -NHS(O)₂R¹⁴, -S(O)₂NHR¹⁵, -C(O)NR¹⁵R¹⁵, -CO₂R¹⁷, -C(O)NR¹⁸R²⁴, -CH₂OH oder 1H- oder 2H-Tetrazol-5-yl steht;

und die pharmazeutisch verträglichen Salze davon,

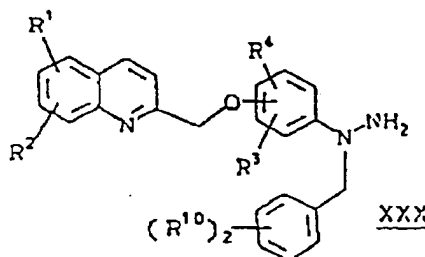
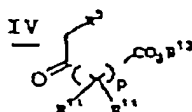
welches umfaßt:

Kuppeln einer Verbindung der Formel XIV mit einer Verbindung der Formel VII



so daß sich eine Verbindung der Formel (I) ergibt, worin (CR¹¹R¹¹)_n-Y_m-(CR¹¹R¹¹)_p-Q für (CR¹¹R¹¹)_p-CO₂Me steht, und danach, falls erwünscht, Überführen der Gruppe (CR¹¹R¹¹)_p-CO₂Me in eine weitere Gruppe der Formel (CR¹¹R¹¹)_n-Y_m-(CR¹¹R¹¹)_p-Q.

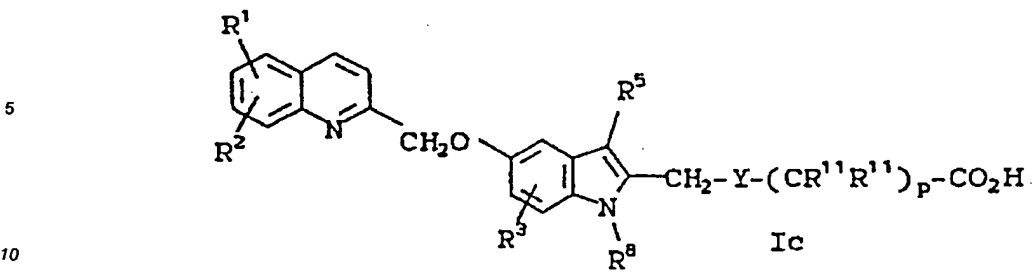
2. Verfahren zur Herstellung der Verbindung der Formel I, worin V für O steht, und R⁸ für CH₂C₆H₃(R¹⁰)₂ steht, wie in Anspruch 1 definiert, welches umfaßt Umsetzen der Verbindung der Formel XXX mit dem Keton der Formel IV durch eine Fischer-Reaktion, um die Verbindung der Formel I bereitzustellen:



worin V für O steht, und R⁸ für CH₂C₆H₃(R¹⁰)₂ steht, und (CR¹¹R¹¹)_n-Y_m-(CR¹¹R¹¹)_p-Q für (CR¹¹R¹¹)_p-CO₂Me steht, und danach, falls erwünscht, Überführen der Gruppe (CR¹¹R¹¹)_p-CO₂Me in eine weitere Gruppe der Formel (CR¹¹R¹¹)_n-Y_m-(CR¹¹R¹¹)_p-Q.

3. Verfahren nach Anspruch 1 zur Herstellung einer Verbindung der Formel (I) nach Anspruch 1, worin n für 0 steht, m für 0 steht, und Q CO₂H bedeutet, oder eines pharmazeutisch verträglichen Salzes davon.
4. Verfahren nach Anspruch 2 zur Herstellung einer Verbindung der Formel (I) nach Anspruch 2, worin n für 0 steht, m für 0 steht, und Q für CO₂H steht, oder eines pharmazeutisch verträglichen Salzes davon.
5. Verfahren nach einem der Ansprüche 1 bis 4 zur Herstellung der Verbindung der Formel Ic oder eines pharmazeutisch verträglichen Salzes davon, worin die Substituenten wie folgt lauten:

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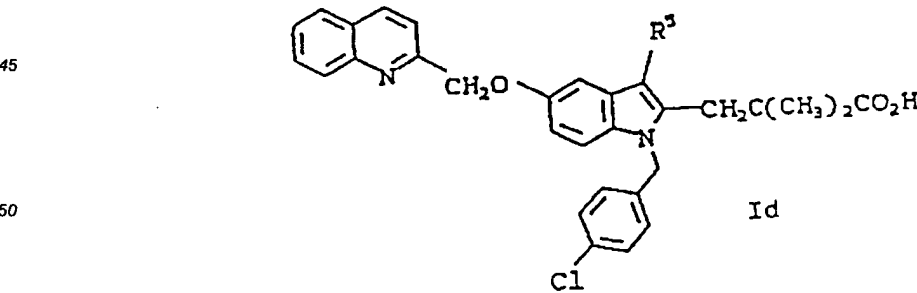
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R ¹	R ²	R ³	R ⁵	R ⁸	Y-(CR ¹¹ R ¹¹) _p
6-Cl	7-Cl	H	Me	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	7-Cl	H	Me	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	4-Allyl	-S-t-Bu	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	4-Allyl	H	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	H	-O-i-Pr	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CH ₂ OCH(Me);
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CHMe;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-S-t-Bu	C(Me) ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CH ₂ OCH ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CHEt;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-Cl	CHMe;
H	H	H	-C(O)CH ₂ -t-Bu	H	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-CF ₃	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-3-OMe	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ CHCH ₂	CH ₂ OCH(Me);
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-OMe	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	Me	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-F	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-Br	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-I	C(Me) ₂ oder
H	H	H	-t-Bu	-CH ₂ Ph-4-F	C(Me) ₂ .

6. Verfahren nach einem der Ansprüche 1 bis 4 zur Herstellung einer Verbindung der Formel Id:



oder eines pharmazeutisch verträglichen Salzes davon, worin R⁵ die in Anspruch 1 gegebene Definition besitzt.

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7. Verfahren nach Anspruch 6 zur Herstellung der Verbindung der Formel Id, worin
 der Substituent R⁵ für folgendes steht: Wasserstoff, -Me, -S-t-Bu, -SPh, -S(O)₂Ph, -S(O)Ph, C(O)Ph, -CH₂Ph, -C(O)CH₂-t-Bu, -O-i-Pr, -C(O)-CF₃, -S(O)₂-t-Bu, S(O)-t-Bu, -C(O)Ph-4-t-Bu, -C(O)Ph-4-Cl, t-Bu, -C(O)Me, -C(O)-c-Pr, -C(O)CH₂CH₂-c-C₅H₉, -C(O)CH₂CH(Me)₂, -C(O)Et, -C(O)CH(Me)₂, -C(O)C(Me)₃,
 5 -C(O)CH₂Ph, -C(Me)₂Pr, -C(Me)₂Et, -CH(Me)₂, -c-Pr, -(i-Me)-c-Pr, -c-C₅H₉, -c-C₆H₁₁, -C(Me)₂Ph, -C-(Me)₂Ph-4-Cl, -1-Ad, -CH₂-1-Ad oder -CH₂CH₂-t-Bu.
8. Verfahren nach einem der Ansprüche 1 bis 4 zur Herstellung einer Verbindung, die folgendermaßen heißt:
- 10 3-[N-(p-Chlorbenzyl)-3-(t-butylthio)-5-chinolin-2-yl-methoxy]-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-methyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Butylthiobenzyl)-3-(t-butylthio)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-(phenylthio)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 15 3-[N-(p-Chlorbenzyl)-3-(phenylsulfonyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure-N-Oxid,
 3-[N-(p-Chlorbenzyl)-3-(phenylsulfonyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-(phenylsulfinyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 20 3-[N-(p-Chlorbenzyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-benzoyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-benzyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 25 2-[N-(p-Chlorbenzyl)-3-(t-butylthio)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-ethoxyethansäure,
 3-[N-(p-Chlorbenzyl)-3-(3,3-dimethyl-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-(t-butylthio)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2-methylpropansäure,
 30 3-[N-(p-Chlorbenzyl)-3-methyl-5-(6,7-dichlorchinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-3-methyl-5-(7-chlorchinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-4-allyl-5-(chinolin-2-yl-methoxy)-3-(t-butylthio)-indol-2-yl]-2,2-dimethylpropansäure,
 35 3-[N-(p-Chlorbenzyl)-4-allyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-6-(chinolin-2-yl-methoxy)-3-(t-butylthio)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-4-(chinolin-2-yl-methoxy)-3-(t-butylthio)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(p-Chlorbenzyl)-7-(chinolin-2-yl-methoxy)-3-(t-butylthio)-indol-2-yl]-2,2-dimethylpropansäure,
 2-[2-[N-(p-Chlorbenzyl)-3-(t-butylthio)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-ethoxy]-propansäure,
 40 3-[N-(p-Chlorbenzyl)-4-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-methyl-3-(p-Chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-methyl-3-(p-Chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Chlorbenzyl)-3-i-propoxy-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Chlorbenzyl)-3-(t-butylthio)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2-ethylpropansäure,
 45 3-[N-(4-Chlorbenzyl)-3-trifluoracetyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Chlorbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2-dimethylpropansäure,
 3-[3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(4-Trifluormethylbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 50 3-[N-Benzyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-(3-Methoxybenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 3-[N-Allyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
 55 3-[N-(4-Methoxybenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,

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- 3-[N-Methyl-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[3-(4-Chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(Phenylsulfonyl)-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-Benzyl-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(t-butylsulfonyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(t-butylsulfinyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-Allyl-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(n-Propyl)-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-Ethyl-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(4-t-butylbenzoyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(4-chlorbenzoyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(1,1-dimethylethyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-acetyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-cyclopropanecarbonyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(3-cyclopentylpropanoyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(3-methylbutanoyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-propanoyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(2-methylpropanoyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-trimethylacetyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-phenylacetyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Fluorbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Brombenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Iodbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(1,1-dimethylbutyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(1,1-dimethylpropyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(3-Fluorbenzyl)-3-(1,1-dimethylethyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(1-methylethyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-cyclopropyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(1-methyl-1-cyclopropyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-cyclopentyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-cyclohexyl-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(α,α -dimethylbenzyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(2-{4-chlor- α,α -dimethylbenzyl})-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-(1-adamantyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(4-Chlorbenzyl)-3-((1-adamantyl)-methyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure,
3-[N-(1,1-Dimethylethyl)-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

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dimethylpropansäure,

3-[N-(1,1-Dimethylpropyl)-3-(4-chlorbenzyl)-6-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

dimethylpropansäure,

3-[N-(4-Chlorbenzyl)-3-(3,3-dimethyl-1-oxo-1-butyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

5 dimethylpropansäure,

Methyl-3-[N-(4-chlorbenzyl)-3,6-bis-(acetyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

dimethylpropanoat oder

Methyl-3-[N-(4-chlorbenzyl)-3,6-bis-(cyclopropancarbonyl)-5-(chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

dimethylpropanoat,

10 oder eines pharmazeutisch verträglichen Salzes davon.

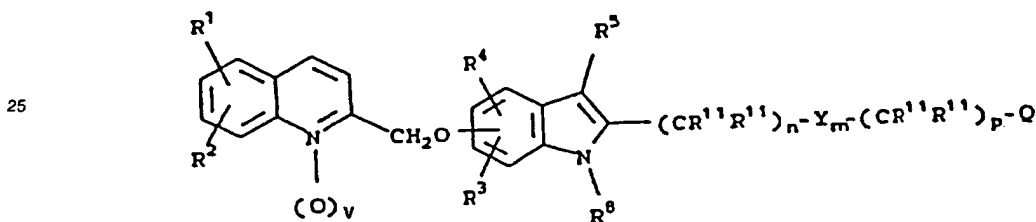
9. Verfahren zur Herstellung von 3-[N-(p-Chlorbenzyl)-3-(t-butylthio)-5-chinolin-2-yl-methoxy)-indol-2-yl]-2,2-dimethylpropansäure oder eines pharmazeutisch verträglichen Salzes davon, welches umfaßt Hydrolyse von Methyl-3-[N-(p-chlorbenzyl)-3-(t-butylthio)-5-chinolin-2-yl-methoxy)-indol-2-yl]-2,2-

15 dimethylpropanoat.

Revendications

Revendications pour les Etats contractants suivants : AT, BE, CH, DE, DK, FR, GB, IT, LI, LU, NL, SE

- 20 1. Composé de formule I :



I

dans laquelle :

R¹, R², R³, R⁴ et R¹⁰ représentent indépendamment un hydrogène, halogène, alkyle C₁₋₇, alcényle C₂₋₇, alcyne C₂₋₇,

35 -CF₃, -CN, -NO₂, -N₃, -C(OH)R¹¹R¹¹ -CO₂R¹² -SR¹⁴, -S(O)R¹⁴, -S(O)₂R¹⁴, -S(O)₂NR¹⁵R¹⁵, -OR¹⁵, -NR¹⁵R¹⁵, -C(O)R¹⁶ ou -(CH₂)_tR²¹;

R⁵ est un hydrogène, -CH₃, -CF₃, -C(O)H, X¹-R⁶ ou X²-R⁷;

R⁶ et R⁹ sont indépendamment : un alkyle, -(CH₂)_uPh(R¹⁰)₂ ou -(CH₂)_uTh(R¹⁰)₂;

40 R⁷ est -CF₃ ou R⁶ ;

R⁸ est un hydrogène ou X³-R⁹ ;

chaque R¹¹ représente indépendamment un hydrogène ou un alkyle C₁₋₇, ou deux R¹¹ sur le même atome de carbone sont réunis pour former un cycle cycloalkyle de 3 à 6 atomes de carbone ;

R¹² est un hydrogène, alkyle C₁₋₇, ou -CH₂R²¹ ;

45 R¹³ est un alkyle C₁₋₇, ou -(CH₂)_tR²¹ ;

R¹⁴ est -CF₃ ou R¹³ ;

R¹⁵ est un hydrogène, -C(O)R¹⁶, R¹³ ou deux R¹⁵, sur le même atome d'azote peuvent être réunis pour former un cycle monocyclique hétérocyclique de 4 à 6 atomes contenant jusqu'à deux hétéroatomes choisis parmi O, S ou N ;

50 R¹⁶ est un hydrogène, -CF₃, alkyle C₁₋₇, alcényle C₂₋₇, alcyne C₂₋₇, ou -(CH₂)_tR²¹ ;

R¹⁷ est -(CH₂)_s-C(R¹⁸R¹⁸)-(CH₂)_s-R¹⁹ ou -CH₂C(O)NR¹⁵R¹⁵;

R¹⁸ est un hydrogène ou un alkyle C₁₋₇ ;

R¹⁹ est

55 a) un cycle hétérocyclique monocyclique ou bicyclique contenant de 3 à 9 atomes de carbone sur le noyau et un ou deux hétéroatomes sur le noyau choisis parmi M, S ou O et chaque cycle dans le radical hétérocyclique est formé de 5 à 6 atomes, ou

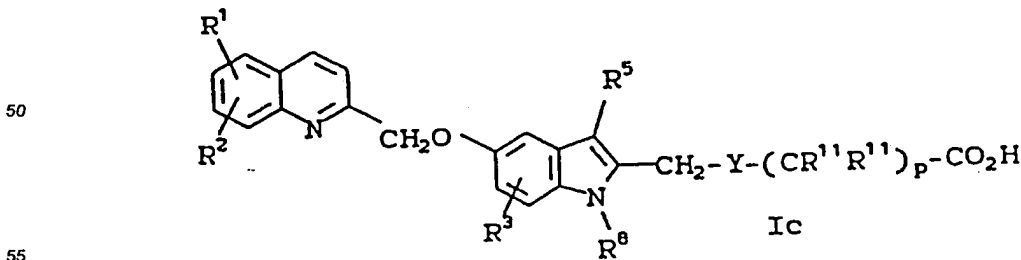
b) le radical W-R²⁰ ;

R²⁰ est un alkyle ou -C(O)R²³ ;

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R²¹ est un phényle substitué avec 1 ou deux groupes R²² ;
 R²² est un hydrogène, un halogène, un alkyle C₁₋₇, alcoxy C₁₋₇, alkylthio C₁₋₇, alkylsulfonyl C₁₋₇,
 alkylcarbonyl C₁₋₇, -CF₃, -CN, -NO₂ ou -N₃ ;
 R²³ est un alkyle, cycloalkyle, ou un cycle monocyclique monohétérocyclique ;
 5 R²⁴ est une structure résiduelle d'acide aminé classique ou R¹⁸ et R²⁴ fixés au même M peuvent être
 cyclisés pour former un reste proline ;
 m est 0 à 1 ;
 n est 0 à 3 ;
 p est 1 à 3 lorsque m est 1 ;
 10 p est 0 à 3 lorsque m est 0 ;
 r est 0 à 2 ;
 s est 0 à 3 ;
 t est 0 à 2 ;
 u est 0 à 3 ;
 15 v est 0 ou 1 ;
 W est O, S ou NR¹⁵,
 X¹ est O ou NR¹⁵,
 X² est C(O), CR¹¹R¹¹, S, S(O) ou S(O)₂ ;
 X³ est C(O), CR¹¹R¹¹, S(O)₂ ou une liaison ;
 20 Y est X¹ ou X² ;
 Q est, -CO₂R¹², -C(O)NHS(O)₂R¹⁴, -NHS(O)₂R¹⁴, -S(O)₂NHR¹⁵, -C(O)NR¹⁵R¹⁵, -CO₂R¹⁷, -C(O)NR¹⁸R²⁴,
 -CH₂OH, ou 1H- ou 2H-tétrazol-5-yl ;
 et leurs sels pharmaceutiquement acceptables.

- 25 2. Composé selon la revendication 1, dans lequel :
 R¹, R², R³ et R⁴ représentent l'hydrogène ;
 R⁵ est X²-R⁷ ou OR⁶ ;
 R⁷ est R⁶ ;
 R⁸ est R⁹ ;
 30 R¹⁰ est un hydrogène comme halogène ;
 m est 0 ;
 n est 1 à 3 ;
 u est 0 dans R⁶ et 1 dans R⁹ ;
 v est 0 ;
 35 X² est CR¹¹ R¹² ou S ;
 Q est -CO₂R¹² ;
 ou un de ses sels pharmaceutiquement acceptables.
3. Composé selon la revendication 1 dans lequel n est 0, m est 0, v est 0 et Q est CO₂H ou son sel
 40 pharmaceutiquement acceptable.
4. Composé selon l'une des revendications 1 à 3, dans lequel R⁵ est SR⁷.
5. Composé selon la revendication 1 de formule Ic, dans lequel les substituants sont les suivants :
- 45

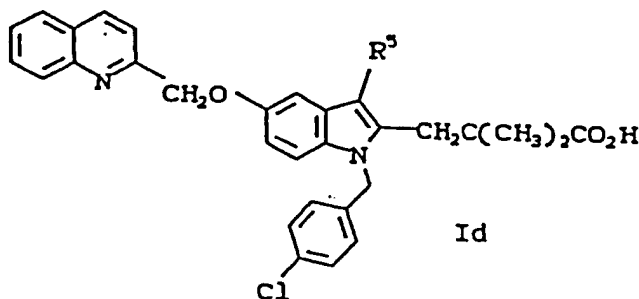


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R ¹	R ²	R ³	R ⁵	R ⁸	Y-(CR ¹¹ R ¹¹) _p
6-Cl	7-Cl	H	Me	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	7-Cl	H	Me	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	4-allyl	-S-t-Bu	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	4-allyl	H	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	H	-O-i-Pr	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CH ₂ OCH(Me);
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CHMe;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-S-t-Bu	C(Me) ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CH ₂ OCH ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CHET;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-Cl	CHMe;
H	H	H	-C(O)CH ₂ -t-Bu	H	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-CF ₃	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-3-OMe	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ CHCH ₂	CH ₂ OCH(Me);
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-OMe	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	Me	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-F	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-Br	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-I	C(Me) ₂ or
H	H	H	-t-Bu	-CH ₂ Ph-4-F	C(Me) ₂ ;

et leurs sels pharmaceutiquement acceptables.

6. Composé selon la revendication 1 de formule Id :



dans laquelle R⁵ est tel que défini à la revendication 1, ou ses sels pharmaceutiquement acceptables.

7. Composé selon la revendication 6, dans lequel le substituant R⁵ est : hydrogène, -Me, -S-t-Bu, -SPh, -S(O)₂Ph, -S(O)Ph, C(O)Ph, -CH₂Ph, -C(O)CH₂-t-Bu, -O-i-Pr, -C(O)-CF₃, -S(O)₂-t-Bu, -S(O)-t-Bu, -C(O)-Ph-4-t-Bu, -C(O)Ph-4-Cl, t-Bu, -C(O)Me, -C(O)-c-Pr, -C(O)CH₂CH₂-c-C₅H₉, -C(O)CH₂CH(Me)₂, -C(O)Et, -C(O)CH(Me)₂, -C(O)C(Me)₃, -C(O)CH₂Ph, -C(Me)₂Pr, -C(Me)₂Et, -CH(Me)₂, -c-Pr, -(l-Me)-c-Pr, -c-C₅H₉, -c-C₆H₁₁, -C(Me)₂Ph, -C(Me)₂Ph-4-Cl, -1-Ad, -CH₂-1-Ad ou -CH₂CH₂-t-Bu.

8. Composé selon la revendication 1, qui est :

Acide 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(p-chlorobenzyl)-3-méthyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(p-t-butylthiobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(p-chlorobenzyl)-3-(phénylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

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Acide 3-[N-(p-chlorobenzyl)-3-(phénylsulfonyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque, N-oxyde ;
 Acide 3-[N-(p-chlorobenzyl)-3-(phénylsulfonyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 5 Acide 3-[N-(p-chlorobenzyl)-3-(phénylsulfonyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-(p-chlorobenzyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-(p-chlorobenzyl)-3-benzoyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 ;
 10 Acide 3-[N-(p-chlorobenzyl)-3-benzyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-(p-chlorobenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-éthoxyéthanoïque ;
 15 Acide 3-[N-(p-chlorobenzyl)-3-(3,3-diméthyl-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-méthylpropanoïque ;
 Acide 3-[N-(p-chlorobenzyl)-3-méthyl-5-(6,7-dichloro-quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 20 Acide 3-[N-(p-chlorobenzyl)-3-méthyl-5-(7-chloro-quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-(p-chlorobenzyl)-4-allyl-5-(quinoléin-2-ylméthoxy)-3-(t-butylthio)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-(p-chlorobenzyl)-4-allyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 25 Acide 3-[N-(p-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)-3-(t-butylthio)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-(p-chlorobenzyl)-4-(quinoléin-2-ylméthoxy)-3-(t-butylthio)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-(p-chlorobenzyl)-7-(quinoléin-2-ylméthoxy)-3-(t-butylthio)indol-2-yl]-2,2-diméthylpropanoïque ;
 30 Acide 2-[2-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-éthoxy]-propanoïque ;
 Acide 3-[N-(p-chlorobenzyl)-4-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-méthyl-3-(p-chlorobenzoyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 35 ;
 Acide 3-[N-méthyl-3-(p-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-(4-chlorobenzyl)-3-i-propoxy-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-(4-chlorobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-éthylpropanoïque ;
 40 ;
 Acide 3-[N-(4-chlorobenzyl)-3-trifluoroacétyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-(4-chlorobenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2-méthylpropanoïque ;
 45 Acide 3-[3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 ;
 Acide 3-[N-(4-trifluorométhylbenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-benzyl-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 50 Acide 3-[N-(3-méthoxybenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-allyl-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 55 Acide 3-[N-(4-méthoxybenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
 Acide 3-[N-méthyl-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

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- Acide 3-[3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(phénylsulfonyl)-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-benzyl-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
5 Acide 3-[N-(4-chlorobenzyl)-3-(t-butylsulfonyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-chlorobenzyl)-3-(t-butylsulfinyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-allyl-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
10 Acide 3-[N-(n-propyl)-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-éthyl-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-chlorobenzyl)-3-(4-t-butylbenzoyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
15 Acide 3-[N-(4-chlorobenzyl)-3-(4-chlorobenzoyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-chlorobenzyl)-3-(1,1-diméthyléthyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-chlorobenzyl)-3-acétyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
20 Acide 3-[N-(4-chlorobenzyl)-3-cyclopropanecarbonyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-chlorobenzyl)-3-(3-cyclopentylpropanoyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-chlorobenzyl)-3-(1-méthyléthyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
25 Acide 3-[N-(4-chlorobenzyl)-3-cyclopropyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-chlorobenzyl)-3-(1-méthyl-1-cyclopropyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
30 Acide 3-[N-(4-chlorobenzyl)-3-cyclopentyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-chlorobenzyl)-3-cyclohexyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-chlorobenzyl)-3-(α,α -diméthylbenzyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
35 Acide 3-[N-(4-chlorobenzyl)-3-(2-(4-chloro- α,α -diméthylbenzyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-chlorobenzyl)-3-(1-adamantyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
40 Acide 3-[N-(1,1-diméthyléthyl)-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(1,1-diméthylpropyl)-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-chlorobenzyl)-3-(3-méthylbutanoyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
45 Acide 3-[N-(4-chlorobenzyl)-3-propanoyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-chlorobenzyl)-3-(2-méthylpropanoyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
50 Acide 3-[N-(4-chlorobenzyl)-3-triméthylacétyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-chlorobenzyl)-3-phénylacétyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-fluorobenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
55 Acide 3-[N-(4-bromobenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
Acide 3-[N-(4-iodobenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-di-

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méthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(1,1-diméthylbutyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(1,1-diméthylpropyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(3-fluorobenzyl)-3-(1,1-diméthyléthyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diéthylpropanoïque ;

3-[N-(4-chlorobenzyl)-3,6-bis(acétyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoate de méthyle

3-[N-(4-chlorobenzyl)-3,6-bis(cyclopropanecarbonyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoate de méthyle,

ou un de leurs sels pharmaceutiquement acceptables.

9. Acide 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ou un de ses sels pharmaceutiquement acceptables.

10. Sel pharmaceutiquement acceptable selon l'une quelconque des revendications 1 à 8, qui est un sel de sodium.

11. Composition pharmaceutique comprenant une quantité efficace d'un composé de la revendication 1 et un véhicule pharmaceutiquement acceptable.

12. Composition pharmaceutique selon la revendication 11, qui comprend en outre une quantité efficace d'un second ingrédient actif choisi dans le groupe constitué des médicaments anti-inflammatoires non-stéroïdiens, des agents analgésiques périphériques, des inhibiteurs de cyclooxygénase, des antagonistes de leucotriène, des inhibiteurs de biosynthèse de leucotriène, des antagonistes de récepteur H₂, des agents anti-histaminiques, des antagonistes de prostaglandine, des antagonistes de thromboxane, des inhibiteurs de synthétase de thromboxane, et des antagonistes ACE.

13. Utilisation d'un composé selon l'une quelconque des revendications 1 à 10 pour préparer un médicament utile à la prévention de la synthèse, de l'action, ou de la sécrétion de SRS-A ou de leucotriènes chez un mammifère.

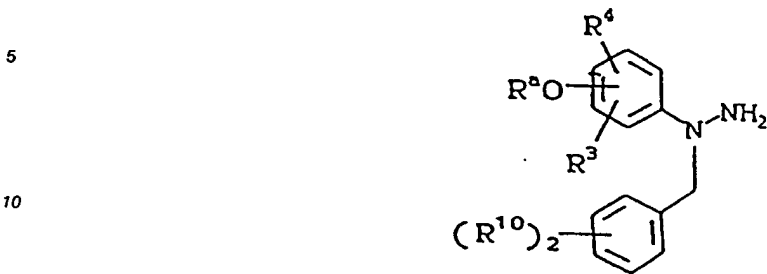
14. Utilisation d'un composé selon l'une quelconque des revendications 1 à 10 pour préparer un médicament utile dans le traitement de l'asthme chez un mammifère.

15. Utilisation d'un composé selon l'une quelconque des revendications 1 à 10 pour préparer un médicament utile au traitement des maladies inflammatoires de l'oeil chez un mammifère.

16. Utilisation d'un composé selon l'une quelconque des revendications 1 à 10 pour préparer un médicament adapté à l'administration d'un composé selon l'une quelconque des revendications 1 à 10, utile au traitement de la maladie inflammatoire de l'intestin chez un patient, au traitement de l'arthrite chez un patient, au traitement de la rhinite allergique chez un patient, au traitement de la conjonctivite allergique chez un patient ou au traitement de l'insuffisance rénale provenant de l'ischémie induite par la toxicologie chimique ou immunologique chez un patient.

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17. Composé de formule :



15 dans laquelle :
 R^a est un t-butylcarbonyle ou



25 R^1 , R^2 , R^3 , R^4 et R^{10} sont tels que définis dans la revendication 1.

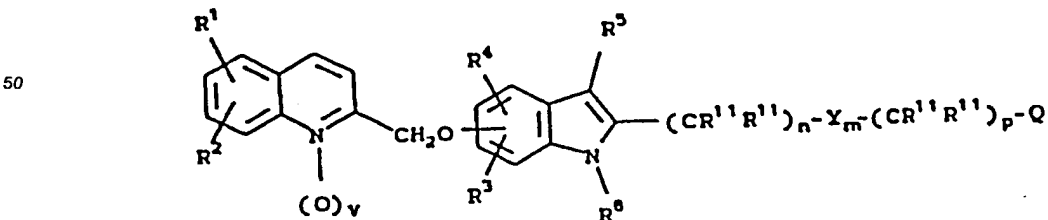
18. Composé selon la revendication 17, dans lequel R^a est



40 19. 1-(p-chlorobenzyl)-1-[4-(quinoléin-2-yl-méthoxy)phényl]hydrazine.

Revendications pour les Etats contractants suivants : ES, GR

45 1. Procédé de préparation d'un composé de formule I :



I

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dans laquelle :

R^1 , R^2 , R^3 , R^4 et R^{10} représentent indépendamment un hydrogène, halogène, alkyle C_{1-7} , alcényle C_{2-7} , alcynyle C_{2-7} ,

$-CF_3$, $-CN$, $-NO_2$, $-N_3$, $-C(OH)R^{11}R^{11}$, $-CO_2R^{12}$, $-SR^{14}$, $-S(O)R^{14}$, $-S(O)_2R^{14}$, $-S(O)_2NR^{15}R^{15}$, $-OR^{15}$,
5 $-NR^{15}R^{15}$, $-C(O)R^{16}$ or $-(CH_2)_lR^{21}$;

R^5 est un hydrogène, $-CH_3$, $-CF_3$, $-C(O)H$, X^1-R^6 or X^2-R^7 ;

R^6 et R^9 sont indépendamment : un alkyle, $-(CH_2)_uPh(R^{10})_2$ ou $-(CH_2)_uTh(R^{10})_2$;

R^7 est $-CF_3$ ou R^6 ;

R^8 est un hydrogène ou X^3-R^9 ;

10 chaque R^{11} représente indépendamment un hydrogène ou un alkyle C_{1-7} , ou deux R^{11} sur le même atome de carbone sont réunis pour former un cycle cycloalkyle de 3 à 6 atomes de carbone ;

R^{12} est un hydrogène, alkyle C_{1-7} , ou $-CH_2R^{21}$;

R^{13} est un alkyle C_{1-7} , ou $-(CH_2)_lR^{21}$;

R^{14} est $-CF_3$ ou R^{13} ;

15 R^{15} est un hydrogène, $-C(O)R^{16}$, R^{13} ou deux R^{15} , sur le même atome d'azote peuvent être réunis pour former un cycle monocyclique hétérocyclique de 4 à 6 atomes contenant jusqu'à deux hétéroatomes choisis parmi O, S ou N ;

R^{16} est un hydrogène, $-CF_3$, alkyle C_{1-7} , alcényle C_{2-7} , alcynyle C_{2-7} , ou $-(CH_2)_lR^{21}$;

R^{17} est ou

20 R^{18} est un hydrogène ou un alkyle C_{1-7} ;

R^{19} est

a) un cycle hétérocyclique monocyclique ou bicyclique contenant de 3 à 9 atomes de carbone sur le noyau et un ou deux hétéroatomes sur le noyau choisis parmi M, S ou O et chaque cycle dans le radical hétérocyclique est formé de 5 à 6 atomes, ou

25 b) le radical $W-R^{20}$;

R^{20} est un alkyle ou $-C(O)R^{23}$;

R^{21} est un phényle substitué avec 1 ou deux groupes R^{22} ;

R^{22} est un hydrogène, un halogène, un alkyle C_{1-7} , alcoxy C_{1-7} , alkylthio C_{1-7} , alkylsulfonyl C_{1-7} , alkylcarbonyl C_{1-7} , $-CF_3$, $-CN$, $-NO_2$ ou $-N_3$;

30 R^{23} est un alkyle, cycloalkyle, ou un cycle monocyclique monohétérocyclique ;

R^{24} est une structure résiduelle d'acide aminé classique ou R^{18} et R^{24} fixés au même N peuvent être cyclisés pour former un reste proline ;

m est 0 à 1 ;

n est 0 à 3 ;

35 p est 1 à 3 lorsque m est 1 ;

p est 0 à 3 lorsque m est 0 ;

r est 0 à 2 ;

s est 0 à 3 ;

t est 0 à 2 ;

40 u est 0 à 3 ;

v est 0 ou 1 ;

W est O, S ou NR^{15} ,

X^1 est O ou NR^{15} ,

X^2 est $C(O)$, $CR^{11}R^{11}$, S, $S(O)$ ou $S(O)_2$;

45 X^3 est $C(O)$, $CR^{11}R^{11}$, $S(O)_2$ ou une liaison ;

Y est X^1 ou X^2 ;

Q est

$-CO_2R^{12}$, $-C(O)NHS(O)_2R^{14}$, $-NHS(O)_2R^{14}$, $-S(O)_2NHR^{15}$, $-C(O)NR^{15}R^{15}$, $-CO_2R^{17}$, $-C(O)NR^{18}R^{24}$,
50 $-CH_2OH$, ou 1H- ou 2H-tétrazol-5-yl ;

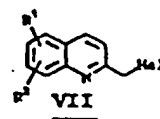
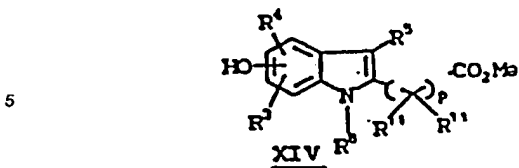
et leurs sels pharmaceutiquement acceptables,

qui comprend :

d'ajouter un composé de formule XIV à un composé de formule VII

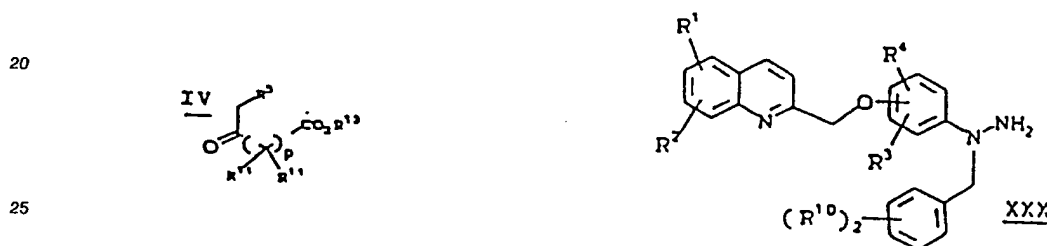
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10 pour donner un composé de formule (I), dans lequel $(CR^{11}R^{11})_n-Y_m-(CR^{11}R^{11})_p-Q$ est $(CR^{11}R^{11})_pCO_2Me$ et ensuite on souhaite transformer le groupe $(CR^{11}R^{11})_pCO_2Me$ en un autre groupe de formule $(CR^{11}R^{11})_n-Y_m-(CR^{11}R^{11})_p-Q$.

2. Procédé de préparation du composé de formule I dans lequel v est 0 et R^8 est $CH_2C_6H_3(R^{10})_2$, tel que c'est défini dans la revendication 1, qui comprend de faire réagir le composé de formule XXX avec la cétone de formule IV par réaction de Fischer pour donner les composés de formule I :

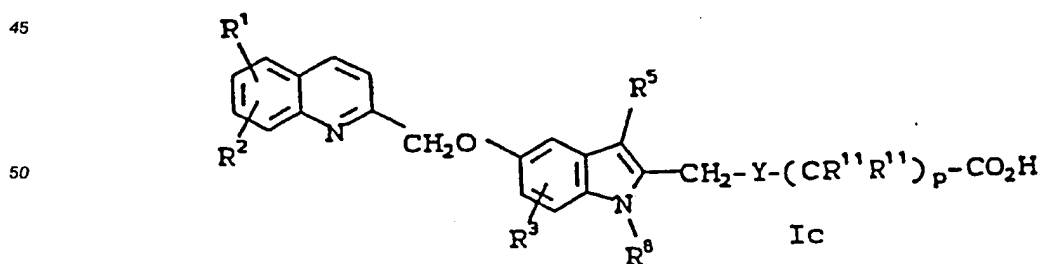


30 dans laquelle v est 0 et R^8 est $CH_2C_6H_3(R^{10})_2$ et $(CR^{11}R^{11})_n-Y_m-(CR^{11}R^{11})_p-Q$ est $(CR^{11}R^{11})_pCO_2Me$ et ensuite, si c'est souhaité, on transforme le groupe $(CR^{11}R^{11})_pCO_2Me$ en un autre groupe de formule $(CR^{11}R^{11})_n-Y_m-(CR^{11}R^{11})_p-Q$.

3. Procédé selon la revendication 1 pour préparer un composé de formule (I) comme c'est défini dans la revendication 1, dans lequel n est 0, m est 0 et Q est CO_2H ou un de ses sels pharmaceutiquement acceptables.

4. Procédé selon la revendication 2 pour préparer un composé de formule (I) tel que défini à la revendication 2 dans lequel n est 0, m est 0, et Q est CO_2H ou un de ses sels pharmaceutiquement acceptables.

5. Procédé selon l'une quelconque des revendications 1 à 4 pour préparer le composé de formule Ic ou un de ses sels pharmaceutiquement acceptables, dans lequel les substituants sont les suivants :

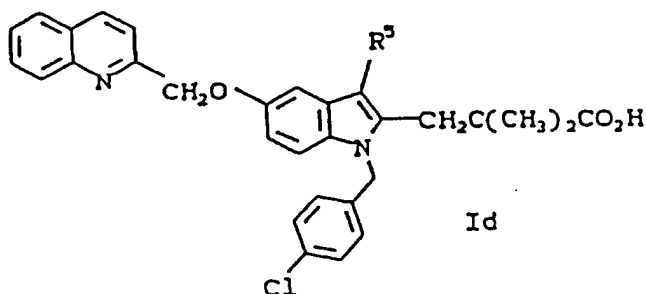


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R ¹	R ²	R ³	R ⁵	R ⁸	Y-(CR ¹¹ R ^{11'}) _p
6-Cl	7-Cl	H	Me	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	7-Cl	H	Me	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	4-allyl	-S-t-Bu	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	4-allyl	H	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	H	-O-i-Pr	-CH ₂ Ph-4-Cl	C(Me) ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CH ₂ OCH(Me);
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CHMe;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-S-t-Bu	C(Me) ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CH ₂ OCH ₂ ;
H	H	H	-S-t-Bu	-CH ₂ Ph-4-Cl	CHEt;
H	H	H	-C(O)CH ₂ -t-	-CH ₂ Ph-4-Cl	CHMe;
H	H	H	-C(O)CH ₂ -t-Bu	H	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-CF ₃	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-3-OMe	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ CHCH ₂	CH ₂ OCH(Me);
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-OMe	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	Me	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-F	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-Br	C(Me) ₂ ;
H	H	H	-C(O)CH ₂ -t-Bu	-CH ₂ Ph-4-I	C(Me) ₂ or
H	H	H	-t-Bu	-CH ₂ Ph-4-F	C(Me) ₂ .

6. Procédé selon l'une quelconque des revendications 1 à 4 pour préparer un composé de formule Id :



ou un de ses sels pharmaceutiquement acceptables dans laquelle R⁵ est tel que défini à la revendication 1.

7. Procédé selon la revendication 6 pour préparer le composé de formule Id dans lequel le substituant R⁵ est : hydrogène, -Me, -S-t-Bu, -SPh, -S(O)₂Ph, -S(O)Ph, C(O)Ph, -CH₂Ph, -C(O)CH₂-t-Bu, -O-i-Pr, -C(O)-CF₃, -S(O)₂-t-Bu, -S(O)-t-Bu, -C(O)Ph-4-t-Bu, -C(O)Ph-4-Cl, t-Bu, -C(O)Me, -C(O)-c-Pr, -C(O)CH₂CH₂-c-C₅H₉, -C(O)CH₂CH(Me)₂, -C(O)Et, -C(O)CH(Me)₂, -C(O)C(Me)₃, -C(O)CH₂Ph, -C(Me)₂Pr, -C(Me)₂Et, -CH(Me)₂, -c-Pr, -(l-Me)-c-Pr, -c-C₅H₉, -c-C₆H₁₁, -C(Me)₂Ph, -C(Me)₂Ph-4-Cl, -1-Ad, -CH₂-1-Ad ou -CH₂CH₂-t-Bu.

8. Procédé selon l'une quelconque des revendications 1 à 4 pour préparer un composé qui est :

Acide 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(p-chlorobenzyl)-3-méthyl-5-(quinoléin-2-ylméthoxy) indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(p-t-butylthiobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

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- Acide 3-[N-(p-chlorobenzyl)-3-(phénylthio)-5-(quinoléin-2-ylméthoxy) indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-(p-chlorobenzyl)-3-(phénylsulfonyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque, N-oxyde ;
- 5 Acide 3-[N-(p-chlorobenzyl)-3-(phénylsulfonyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-(p-chlorobenzyl)-3-(phénylsulfinyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-(p-chlorobenzyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- 10 Acide 3-[N-(p-chlorobenzyl)-3-benzoyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-(p-chlorobenzyl)-3-benzyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-(p-chlorobenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- 15 Acide 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-éthoxyéthanoïque ;
- Acide 3-[N-(p-chlorobenzyl)-3-(3,3-diméthyl-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-méthylpropanoïque ;
- 20 Acide 3-[N-(p-chlorobenzyl)-3-méthyl-5-(6,7-dichloro-quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-(p-chlorobenzyl)-3-méthyl-5-(7-chloro-quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-(p-chlorobenzyl)-4-allyl-5-(quinoléin-2-ylméthoxy)-3-(t-butylthio)indol-2-yl]-2,2-diméthylpropanoïque ;
- 25 Acide 3-[N-(p-chlorobenzyl)-4-allyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-(p-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)-3-(t-butylthio)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-(p-chlorobenzyl)-4-(quinoléin-2-ylméthoxy)-3-(t-butylthio)indol-2-yl]-2,2-diméthylpropanoïque ;
- 30 Acide 3-[N-(p-chlorobenzyl)-7-(quinoléin-2-ylméthoxy)-3-(t-butylthio)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 2-[2-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-éthoxy]-propanoïque ;
- 35 Acide 3-[N-(p-chlorobenzyl)-4-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-méthyl-3-(p-chlorobenzoyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-méthyl-3-(p-chlorobenzyl)-6-(quinoléin-2-ylméthoxy) indol-2-yl]-2,2-diméthylpropanoïque ;
- 40 Acide 3-[N-(4-chlorobenzyl)-3-i-propoxy-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-(4-chlorobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-éthylpropanoïque ;
- Acide 3-[N-(4-chlorobenzyl)-3-trifluoroacétyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- 45 Acide 3-[N-(4-chlorobenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2-méthylpropanoïque ;
- Acide 3-[3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- 50 Acide 3-[N-(4-trifluorométhylbenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-benzyl-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-(3-méthoxybenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- 55 2,2-diméthylpropanoïque ;
- Acide 3-[N-allyl-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;
- Acide 3-[N-(4-méthoxybenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-

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2,2-diméthylpropanoïque;

Acide 3-[N-méthyl-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

5 Acide 3-[N-(phénylsulfonyl)-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-benzyl-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(t-butylsulfonyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

10 Acide 3-[N-(4-chlorobenzyl)-3-(t-butylsulfinyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-allyl-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(n-propyl)-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

15 Acide 3-[N-éthyl-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(4-t-butylbenzoyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(4-chlorobenzoyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

20 Acide 3-[N-(4-chlorobenzyl)-3-(1,1-diméthyléthyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-acétyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-cyclopropanecarbonyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

25 Acide 3-[N-(4-chlorobenzyl)-3-(3-cyclopentylpropanoyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(1-méthyléthyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

30 Acide 3-[N-(4-chlorobenzyl)-3-cyclopropyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(1-méthyl-1-cyclopropyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-cyclopentyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

35 Acide 3-[N-(4-chlorobenzyl)-3-cyclohexyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(α,α -diméthylbenzyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

40 Acide 3-[N-(4-chlorobenzyl)-3-(2-(4-chloro- α,α -diméthylbenzyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(1-adamantyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(1,1-diméthyléthyl)-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

45 Acide 3-[N-(1,1-diméthylpropyl)-3-(4-chlorobenzyl)-6-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(3-méthylbutanoyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

50 Acide 3-[N-(4-chlorobenzyl)-3-propanoyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(2-méthylpropanoyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-triméthylacétyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

55 Acide 3-[N-(4-chlorobenzyl)-3-phénylacétyl-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-fluorobenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

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Acide 3-[N-(4-bromobenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-iodobenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

5 Acide 3-[N-(4-chlorobenzyl)-3-(1,1-diméthylbutyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(1,1-diméthylpropyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

10 Acide 3-[N-(3-fluorobenzyl)-3-(1,1-diméthyléthyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ;

Acide 3-[N-(4-chlorobenzyl)-3-(3,3-diméthyl-1-oxo-1-butyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diéthylpropanoïque ;

3-[N-(4-chlorobenzyl)-3,6-bis(acétyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoate de méthyle

15 3-[N-(4-chlorobenzyl)-3,6-bis(cyclopropanecarbonyl)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoate de méthyle,
ou un de leurs sels pharmaceutiquement acceptables.

9. Procédé de préparation de l'acide 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoïque ou un de ses sels pharmaceutiquement acceptables, qui comprend l'hydrolyse du 3-[N-(p-chlorobenzyl)-3-(t-butylthio)-5-(quinoléin-2-ylméthoxy)indol-2-yl]-2,2-diméthylpropanoate de méthyle.

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